

53

/A COMPUTER PROGRAM FOR INTERNAL DOSIMETRY ANALYSIS
USING THE METHODS OF ICRP-30/

by

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TABLE OF CONTENTS

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	<u>page</u>
TABLE OF CONTENTS.	i
LIST OF FIGURES.	iii
LIST OF TABLES	iv
0.0 INTRODUCTION.	1
1.0 FUNDAMENTALS OF RADIATION PROTECTION.	3
1.1 Dose Equivalent H.	4
1.2 Absorbed Dose D.	4
1.3 Effective Quality Factor \bar{Q}	5
1.4 Collective Dose Equivalent S	5
1.4.1 Radiobiological Assumption.	7
1.5 Committed Dose Equivalent H_{50}	7
1.5.1 Non-homogeneous radiation	8
1.5.2 Unequal sensitivity of cells.	9
1.5.3 Rate of dose accumulation	9
1.5.4 Age, sex, and dose-dependency	9
1.6 Dose Equivalent Limits for Occupational Exposure	10
1.6.1 Recommended limit for non-stochastic effects.	10
1.6.2 Recommended limit for stochastic effects.	10
1.6.3 External exposures to penetrating radiation	11
1.6.4 Planned special exposures	12
1.6.5 Women of reproductive capacity.	12
1.6.6 Occupational exposure of pregnant women	12
1.7 Control of Internal Dose	12
1.7.1 Annual limit on intake (ALI).	12
1.7.2 Derived air concentration (DAC)	13
2.0 EVALUATION OF SPECIFIC COMMITTED DOSE EQUIVALENT.	15
2.1 Specific Effective Energy.	18
2.1.1 Target organs of the GI tract and bladder wall.	22
2.1.2 Target organ in bone.	23
2.1.3 A note on daughters	25
2.2 Number of Transformations in a Source Organ over 50 years.	25
2.2.1 A note on daughters	26
2.2.2 Transfer compartment T.	26
2.2.3 Tissue compartment p.	32
2.2.4 Respiratory system.	34
2.2.5 Gastrointestinal tract.	47

	<u>Page</u>
2.2.6 Activity translocated to the transfer compartment f_T^i	54
2.2.7 Source organ as bone.	56
2.2.8 Three compartments model for iodine	57
3.0 DOCUMENTATION FOR "DOSE FORTRAN".	61
3.1 Objective.	61
3.2 Requirements for Execution	62
3.3 Execution of the Program	64
3.4 Dose Fortran	65
3.4.1 Decay1 Fortran	72
3.4.2 Atomno Fortran	77
3.4.3 Iclass Fortran	78
3.4.4 Pelass Fortran	78
3.4.5 Flvalu Fortran	78
3.4.6 Frac Fortran	79
3.4.7 Factor Fortran	81
3.4.8 Icrp Fortran	83
3.4.9 Inhale Fortran	84
3.4.10 Respir Fortran	92
3.4.11 Thalf Fortran.	94
3.4.12 Refman Fortran	95
3.4.13 Tfrac Fortran.	97
3.4.14 Trnsfm Fortran	100
3.4.15 Speff Fortran.	104
3.4.16 Decay Fortran.	108
3.4.17 Yerror Fortran	112
3.4.18 Intrpt Fortran	113
3.4.19 Ingest Fortran	115
3.4.20 Result Fortran	120
3.4.21 Submer Fortran	123
3.5 Results and Discussion	124
4.0 REFERENCES.	145
ACKNOWLEDGEMENTS	146
APPENDIX A: Listing of Computer Programs.	147

LIST OF FIGURES

<u>Figure</u>		<u>Page</u>
1.1	Relationship of units.	6
2.1	Mathematical model used by the ICRP to describe the kinetics of radionuclides in the body.	27
2.2	Mathematical model used to describe clearance from respiratory system.	36
2.3	Model of deposition of dust in the respiratory system as proposed by the ICRP	38
2.4	Pulmonary clearance classification of inorganic compounds as proposed by the ICRP.	39
2.5	Model of gastrointestinal tract as proposed by the ICRP	48
2.6	Mathematical model proposed by the ICRP for radioactive iodine	57
3.1	Points of call of subroutines in execution of the program	66
3.2	Flowchart of "INHALE FORTRAN".	87
3.3	Flowchart of subroutine "TRAC FORTRAN"	99
3.4	Flowchart of "TRANSFM FORTRAN".	106
3.5	Flowchart of function subprogram "SPEFF FORTRAN".	109
3.6	Flow diagram for "INCEST FORTRAN"	119

LIST OF TABLES

<u>Table</u>		<u>Page</u>
1.1	Weighting factors as proposed by the ICRP.	11
2.1	Absorbed fractions for dosimetry of radionuclides in bone as recommended by the ICRP 30.	24
2.2	The values of removal half times and compartmental fractions in the four respiratory regions.	40
2.3	Approximate expressions for the number of transformations in the subcompartments of lung following inhalation of unit Bq of activity.	42
2.4	Approximate expressions for the number of transformations of a radioactive daughter in the subcompartments of the lung. A_a to A_j refers to the number of transformations of the immediate parent in subcompartments a to j of the lung. λ'_R is the radiological decay constant of the daughter	43
2.5	Mathematical model used to describe the kinetics of radionuclides in the gastrointestinal tract as proposed by the ICRP	49
2.6	Approximate expressions for the number of transformations of the parent in the various regions of the gastrointestinal tract following ingestion of 1 Bq of activity. Based on ICRP Publication 30	51
2.7	Approximate expressions for the number of transformations of a radioactive daughter in the various sections of the GI tract. A_{ST} , A_{SI} , A_{ULI} , A_{LLI} are the number of transformations of the parent in the various regions of the tract. λ'_R is the radiological decay constant of the daughter. Based on ICRP Publication 30	52
3.1	Masses of organs and tissues of "Reference Man" in this program.	62
3.2	Specific effective energy absorbed in a target organ from each transformation in a source organ for radionuclide I-131	129

<u>Table</u>	<u>Page</u>
3.3	Specific effective energy absorbed in a target organ from each transformation in a source organ for radionuclide In-113m 130
3.4	Specific effective energy absorbed in a target organ from each transformation in a source organ for radionuclide Te-121. 131
3.5	Specific committed dose equivalents (Sv/Bq) for selected ingested radionuclides and their daughters. Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks. 132
3.6	Specific committed dose equivalents (Sv/Bq) for selected ingested radionuclides and their daughters. Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks. 133
3.7	Specific committed dose equivalents (Sv/Bq) for selected ingested radionuclides and their daughters. Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks. 134
3.8	Specific committed dose equivalents (Sv/Bq) for selected inhaled radionuclides and their daughters. Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks. 135
3.9	Specific committed dose equivalents (Sv/Bq) for selected inhaled radionuclides and their daughters. Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks. 136
3.10	Specific committed dose equivalents (Sv/Bq) for selected inhaled radionuclides and their daughters. Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks. 137
3.11	Specific committed dose equivalents (Sv/Bq) for selected inhaled radionuclides and their daughters. Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks. 138
3.12	Specific committed dose equivalents (Sv/Bq) for selected inhaled radionuclides and their daughters. Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks. 139

<u>Table</u>	<u>Page</u>
3.13	Specific committed dose equivalents (Sv/Bq) for selected inhaled radionuclides and their daughters. Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks. 140
3.14	Comparison of specific committed dose equivalents (Sb/Bq) to the results in ICRP Publication 30 for selected ingested radionuclides and their daughters. Values reported are greater than or equal to 10% of the maximum 141
3.15	Comparison of specific committed dose equivalents (Sb/Bq) to the results in ICRP Publication 30 for selected ingested radionuclides and their daughters. Values reported are greater than or equal to 10% of the maximum 142
3.16	Comparison of specific committed dose equivalents (Sv/Bq) to the results in ICRP Publication 30 for selected inhaled radionuclides and their daughters. Values reported are greater than or equal to 10% of the maximum. 143
3.17	Comparison of specific committed dose equivalents (Sv/Bq) to the results in ICRP Publication 30 for selected inhaled radionuclides and their daughters. Values reported are greater than or equal to 10% of the maximum. 144

0.0 INTRODUCTION

Since the discovery of radioactivity, mankind has been confronted with the challenge of understanding biological effects of radiation. The domestication of nuclear power has heightened the need to fully comprehend the relation between radiation and human health for the safety of occupational workers and the public in general. During recent decades, the field of radiation protection has made tremendous strides from its infancy. The rapid growth of knowledge has made it possible to revise the mathematical models for accurate estimation of internal doses from radionuclides, and update the regulatory guidelines for control and management of radioactive materials. In view of this, the International Commission on Radiological Protection approved a new set of basic recommendations dealing with protection from ionizing radiation. These are detailed in the ICRP Publication 26 [IC77]. Further, the methods used to evaluate internal dose to comply with the limits in ICRP-26 along with the tables of specific committed dose equivalents, limits on annual intake, and air concentrations from specific radionuclides are shown in ICRP Publication 30 [IC79].

This report describes a computer program "DOSE" in FORTRAN-77 that generates internal dose factors of the type called for in ICRP-30. The constraints imposed by the ICRP on exposure of different organs and tissues of the body singly or together with other organs, in the case of intake of a radionuclide, are discussed in Chapter 1.0. The general principles, mathematical models, definitions, and calculational

procedures adopted by the ICRP-30 for calculation of specific committed dose equivalent are used in this program. Also, the models presented in Chapter 2.0 for the routes of entry of radionuclides into the body, namely, the respiratory system and the gastrointestinal tract, are the ones recommended by the ICRP-30. The metabolic data for the radioisotopes of different elements are also obtained from ICRP-30. In the case of iodine, the program follows the alternate model discussed in the ICRP-30. The specific committed dose equivalents are assumed to be received by a "reference man" of anatomical and physiological characteristics described in ICRP Publication [IC75]. The decay schemes of radionuclides are obtained from Kocher [K081]. The ICRP uses the decay schemes from its publication 38 [IC83]. This difference may reflect on the results for some of the radionuclides.

An important feature of this program is its flexibility. The data files on radiological and biological decay are written in such a manner that addition of new data on a radionuclide or amendment of the present data is easy. Also, as is explained in Chapter 3.0, most of the calculations for important quantities are done by separate, independent subroutines. This design makes future modifications in the program convenient and simple.

1.0 FUNDAMENTALS OF RADIATION PROTECTION

The concept of radiation protection centers around two possible effects in individuals. They are defined as "somatic" and "hereditary".

"Somatic" effects of radiation are limited to the exposed individual while "hereditary" effects are manifested in the individual's progeny. The quality of effect can be described as "stochastic" or "non-stochastic".

"Stochastic" effects involve probability of occurrence as opposed to severity, and are therefore considered a function of dose, without threshold. In contrast, "non-stochastic" effects vary in severity with dose, and hence may involve a threshold. Hereditary effects are considered stochastic at the dose range involved in radiation protection, while somatic effects may be stochastic or non-stochastic. An example of stochastic effects is carcinogenesis at low doses. Non-stochastic somatic effects may range from cataract of the lens to damage of the hematopoietic system.

To quantify health damage, the International Commission on Radiological Protection (ICRP) has proposed the concept of "detriment". "Detriment" in a population is described as the mathematical "expectation" of the deleterious effect from a radiation exposure, taking into account not only the probability of each type of damaging effect but also the severity of the effect. In other words, "detriment" to health G in a group of persons P is given as the summation over all effects i of the product of probability p_i of

suffering the effect and the severity of effect expressed by a weighting factor g_i , i.e.,

$$G = P \sum_i p_i g_i . \quad (1.1)$$

1.1 DOSE EQUIVALENT, H

This quantity characterizes the severity and probability of the deleterious health effects of radiation. The dose equivalent H at a point in tissue is given by

$$H = D \bar{Q} N , \quad (1.2)$$

where D is the absorbed dose,

\bar{Q} is the effective quality factor, and

N is the product of all modifying factors such as absorbed dose rate and fractionation specified by ICRP. At present, it is assigned a value of 1.

The SI unit of dose equivalent is the sievert (Sv). $1 \text{ Sv} \equiv 100 \text{ rem}$.

1.2 ABSORBED DOSE, D

This refers to the energy locally deposited (and ultimately dissipated as heat) per unit mass of the medium. It applies to both types of radiation, ionizing and non-ionizing. The SI unit of absorbed dose is the gray (Gy). $1 \text{ Gy} = 1 \text{ J kg}^{-1}$ ($\equiv 100 \text{ rad}$).

1.3 EFFECTIVE QUALITY FACTOR, \bar{Q}

Quality factor is introduced to allow for the effect on the detriment of the microscopic distribution of absorbed energy. It is a function of the collisional stopping power in water. For a spectrum of radiation, the ICRP [IC77] recommends an effective value \bar{Q} to be used for both external and internal radiation:

X rays, γ rays, and electrons	1
Neutrons, protons and singly-charged particles of rest mass greater than one atomic mass unit of unknown energy	10
α particles and multiply-charged particles (and particles of unknown charge) of unknown energy	20

A graphic review of the relationship between these quantities is shown in Fig. 1.1.

1.4 COLLECTIVE DOSE EQUIVALENT, S

This concept approximates the relationship between detriment and the distribution of dose equivalent in an exposed population. It is expressed as:

$$S = \sum_i H_i P_i , \quad (1.3)$$

where H_i is the dose equivalent to the whole body or a specific organ or tissue of an individual who is a member of the subgroup i in the exposed population, and

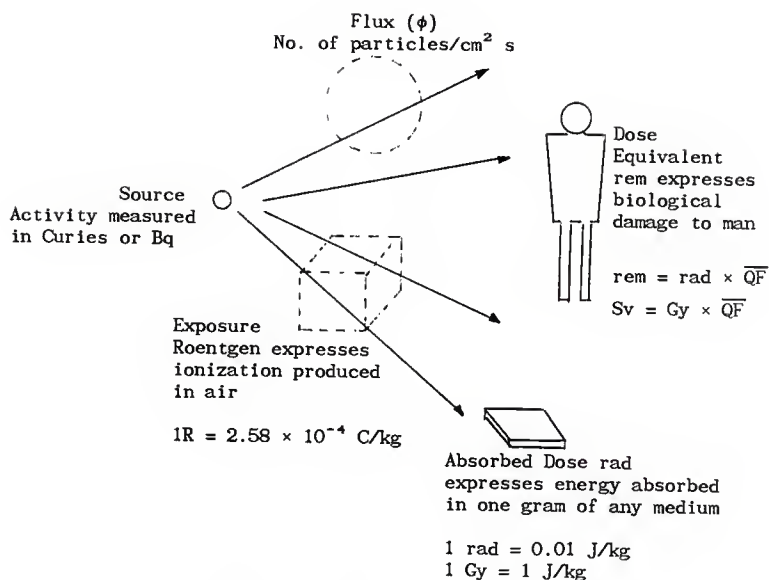


FIG. 1.1. Relationship of units.

P_i represents the number of members in this subgroup i .

The collective dose equivalent S_k from a practice or source (k) is given by

$$S_k = \int_0^{\infty} HP(H)dH , \quad (1.4)$$

where $P(H)dH$ is the number of individuals receiving a dose equivalent in the whole body or a specific organ or tissue in the range H to $H + dH$.

1.4.1 Radiobiological Assumption:

A simple summation over all subgroups of the population of doses to specific organs or tissues of a typical individual in a subgroup is used as a measure of detriment to estimate the collective dose equivalent in a population [IC77]. This process is based on an assumption regarding stochastic effects; namely, that within the usual range of radiation exposure a linear relationship without threshold exists between dose and the probability of an effect.

1.5 COMMITTED DOSE EQUIVALENT, H_{50}

This is defined as the total dose equivalent averaged throughout a tissue in the 50 years after intake of a radionuclide into the body. The 50 year period represents a working life. Mathematically, it can be expressed as:

$$H_{50} = \int_{t_0}^{t_0 + 50y} \dot{H}(t) dt , \quad (1.5)$$

where $\dot{H}(t)$ is the relevant dose-equivalent rate, and t_0 is the time of intake.

In view of the radiobiological assumption above, it is considered practical to conceptualize the term "dose equivalent" as a mean dose equivalent over all cells of uniform sensitivity in a particular tissue or organ for stochastic effects.

1.5.1 Non-homogeneous radiation:

If the doses to individual cells vary due to non-uniform irradiation then the relevance of the mean dose equivalent may be questioned. However, according to the Commission, based on theoretical concepts and epidemiological evidence, a series of "hot spots" in an organ or tissue is less likely to be damaging than a uniform distribution. High doses cause loss of reproductive capacity or death of cells, neither of which may contribute significantly to stochastic effects. Thus, for non-homogeneous irradiation, assessment of dose with the assumption of homogeneous distribution would probably be an overestimation. So far as non-stochastic effects are concerned, the limited amount of cell killing at moderate dose levels is thought not to be a major factor in perturbing the balance of organ function.

1.5.2 Unequal sensitivity of cells:

The problem of dose variance due to differing cell sensitivities of different organs is handled by the Commission by prescribing risk factors according to relative radiosensitivities of the irradiated tissues. A detailed explanation of these risk factors will follow later.

1.5.3 Rate of dose accumulation:

Based on assumptions and explanations summarized above, the Commission has found it unnecessary to recommend a maximum rate of dose accumulation as long as the annual dose equivalent limit is met. The only exception is the case of occupational exposure of women of reproductive potential and pregnant women.

1.5.4 Age, sex, and dose-dependency:

Risk factors for occurrence of malignant cancers are thought to be lessened in older persons due to long periods of latency needed for progression and expression of these effects. Also, development of breast cancer is much more common in females than in males. Such facts emphasize the importance of age, sex, and dose dependency. The Commission however, describes average risk levels based on anatomical and physiological characteristics of "reference man" because, for protection purposes, it is considered sufficiently prudent to use a single dose-equivalent limit for each organ or tissue for all workers regardless of age or sex.

1.6 DOSE EQUIVALENT LIMITS FOR OCCUPATIONAL EXPOSURE

The primary considerations of the Commission in establishing dose equivalent limits were:

- i) For stochastic effects, a single value should not only limit uniform irradiation of the whole body but also ensure that the total risk from irradiation to parts of the body does not exceed the value.
- ii) To preclude non-stochastic effects, a value should be proposed such that no single tissue receives more dose-equivalent than the value.

1.6.1 Recommended limit for non-stochastic effects:

To prevent non-stochastic effects in all tissues except the lens, the Commission has proposed a limit of 0.5 Sv (50 rem) in a year. The dose equivalent to the lens is limited to 0.15 Sv (15 rem) in a year.

1.6.2 Recommended limit for stochastic effects:

For whole-body uniform irradiation, the limit is

$$H_{wb} = 5 \text{ cSv (5 rem) in any year.} \quad (1.6)$$

Non-uniform irradiation must adhere to the following condition:

$$\sum_T w_T H_T \leq H_{wb} \quad (1.7)$$

where H_T is the dose equivalent received by tissue T, and

w_T is a weighting factor involving relative radiosensitivity of the tissue T.

The values of these weighting factors are given in Table 1.1.

TABLE 1.1. Weighting factors as proposed by the ICRP Publication 26.

Tissue	w_T
Gonads	0.25
Breast	0.15
Red bone marrow	0.12
Lung	0.12
Thyroid	0.03
Bone surfaces	0.03
Remainder	0.30

The "remainder" refers to five remaining organs or tissues receiving the highest dose equivalents, with $w_T = 0.06$ applied to each. All other tissues are neglected. Of course, lens and skin are not members of "remainder" because they are subject to non-stochastic effects. The gastrointestinal tract is treated as four separate organs: stomach, small intestine, upper large intestine, and lower large intestine.

1.6.3 External exposures to penetrating radiation:

In the absence of information on actual distribution of dose equivalent within the body, the maximum value of dose equivalent in a 30-cm spherical phantom when limited to the 0.5 Sv annual limit will permit a comparable level of protection.

1.6.4 Planned special exposures:

The limit under such situations should not exceed twice the relevant annual limit in any single event, and, in a lifetime, five times this annual limit. The exposures however, must be justified and permitted only when alternative techniques are impractical or unavailable.

1.6.5 Women of reproductive capacity:

Occupational exposure of such women should be restricted to insure that the embryo receives no more than 0.005 Sv during the first two months of pregnancy.

1.6.6 Occupational exposure of pregnant women:

Exposure should be minimized and not exceed 30% of the normal annual limits.

1.7 CONTROL OF INTERNAL DOSE

For control of internal dose for workers, the standards are derived from the general limits described in the previous section and are based on the parameters of the "reference man".

1.7.1 Annual limit on intake (ALI) [IC79]:

For a given radionuclide, if I is the annual intake (Bq) either by ingestion or inhalation, and $\hat{H}_{50,T}$ (Sv/Bq) is the specific (per unit intake) committed dose equivalent in tissue T from the intake by the

specified mode, weighted for its sensitivity by weighting factor w_T described in Table 1.1, then the annual limit on intake is defined as the greater value of I which satisfies both of the following inequalities

$$I \sum_T w_T \hat{H}_{50,T} \leq 0.05 \text{ Sv} \quad (1.8)$$

for stochastic effects, and

$$I \hat{H}_{50,T} \leq 0.5 \text{ Sv} \quad (1.9)$$

for non-stochastic effects.

The summation in the first inequality is over all tissues in Table 1.1, including of course the five "remaining tissues" in "remainder" which receive the greatest specific committed dose equivalents. A tissue is considered to be "significantly irradiated" only when it satisfies the following inequality

$$w_T \hat{H}_{50,T} \geq 0.1 (w_T \hat{H}_{50,T})_{\text{maximum}} \quad (1.10)$$

1.7.2 Derived air concentration (DAC) [IC79]:

The DAC for any radionuclide is defined as that concentration in air (Bq/m^3) which, if received by "reference man" for a working year of 2000 h (50 weeks at 40 hours per week), would lead to the ALI for inhalation, i.e.,

$$\text{DAC} = \text{ALI}/(2000 \times 60 \times 0.02),$$

or

$$\text{DAC} = \text{ALI}/2.4 \times 10^3 \text{ Bq/m}^3, \quad (1.11)$$

where 0.02 m^3 is the volume of air breathed at work by "reference man" per minute under conditions of "light activity" as described by ICRP Publication 23.

For radioactive noble gases other than radon and thoron, DAC's are the concentrations which would lead to a cumulative weighted average dose of not more than 0.05 Sv, a dose to the lens of 0.15 Sv, or a dose to the skin of 0.5 Sv in a working year (2000 h).

2.0 EVALUATION OF SPECIFIC COMMITTED DOSE EQUIVALENT

Specific committed dose equivalent from a radionuclide to a target organ or tissue is defined as the committed dose equivalent to the organ or tissue as a result of ingestion or inhalation of unit activity of the nuclide.

Activity or decay rate A is mathematically expressed as

$$A = \lim_{\Delta t \rightarrow 0} \frac{\Delta N}{\Delta t}, \quad (2.1)$$

where ΔN is the number of spontaneous nuclear transformations in a quantity of radionuclide, and

Δt is the time interval.

The unit of activity is the becquerel (Bq). $1 \text{ Bq} = 1 \text{ s}^{-1}$ ($\approx 2.7 \times 10^{-11} \text{ Ci}$).

Committed dose equivalent, as explained in the previous chapter, is the total dose equivalent averaged throughout any tissue in the 50 years following intake of a radionuclide into the body, and hence can be written as

$$H_{50} = \sum_i \frac{\int_0^M D_{50,i} \bar{Q}_i N_i dm}{\int_0^M dm}, \quad (2.2)$$

where i is an index for the type of radiation (α, β, γ , etc.) released by the radionuclide, ;

$D_{50,i}$ is the total absorbed dose in the element of mass dm of the specific organ or tissue during a 50 year period following intake of the radionuclide into the body,

\bar{Q}_i is the effective quality factor for the radiation type i ,

N_i is the product of all modifying factors such as dose rate, fractionation, etc., and

M is the mass of the organ or tissue under consideration.

A value of 1 is recommended for N by the ICRP (Publication 26) and the value of \bar{Q} is constant for each type of radiation i ; therefore, the above relation can be simplified to

$$H_{50} = \sum_i \bar{Q}_i \overline{D}_{50,i} \quad (2.3)$$

where $\overline{D}_{50,i}$ is the total absorbed dose during the 50 years after intake of the radionuclide into the body, averaged throughout the specified organ or tissue for each radiation type i .

When a radionuclide enters the body, it is distributed in various source organs or tissues where it may decay or be eliminated by normal metabolic processes. The committed dose equivalent in a target organ or tissue T due to the disintegration of a radionuclide j , releasing radiation of type i in source organ S , denoted by, $H_{50}(T \leftarrow S)_{i,j}$, is defined as a product of two factors:

- i) the total number of transformations (decays) of radionuclide j in source organ S during the 50 year period following its entry into the body, and
- ii) the energy absorbed per unit mass in the target organ T , modified for effective quality factor, from radiation of type i per transformation of radionuclide j in source organ S .

Mathematically, for each radiation type i from radionuclide j ,

$$H_{50}(T \leftarrow S)_{i,j} = \overline{Q}_i \overline{D_{50}}(T \leftarrow S)_{i,j} ,$$

or

$$H_{50}(T \leftarrow S)_{i,j} = U_{s,j} \times 1.6 \times 10^{-13} \times SEE(T \leftarrow S)_{i,j} \times 10^3 , \quad (2.4)$$

where

$U_{s,j}$ is the number of transformations (decays) of radionuclide j in organ S over 50 years following intake of the radionuclide,

1.6×10^{-13} is the number of joules in 1 MeV,

$SEE(T \leftarrow S)_{i,j}$ is the specific effective energy, modified by quality factor, for radiation type i absorbed in T from each transformation in S . The unit is MeV/g per transformation, and

10^3 is the conversion factor from g^{-1} to kg^{-1} .

Thus, for all types of radiation emitted by radionuclide j ,

$$H_{50}(T \leftarrow S)_j = 1.6 \times 10^{-10} \left[U_{s,j} \sum_i SEE(T \leftarrow S)_i \right]_j . \quad (2.5)$$

If the radionuclide decays into a radioactive daughter j' , then

$$H_{50}(T \leftarrow S)_{j+j'} = 1.6 \times 10^{-10} \left\{ \left[U_{s,j} \sum_i SEE(T \leftarrow S)_i \right]_j + \left[U_{s,j'} \sum_i SEE(T \leftarrow S)_i \right]_{j'} \right\} . \quad (2.6)$$

For a number of radioactive daughters, the committed dose equivalent in

target T from decays in source S can be generalized as

$$\sum_j H_{50}(T \leftarrow S)_j = 1.6 \times 10^{-10} \sum_j \left[U_{s,j} \sum_i SEE(T \leftarrow S)_i \right]_j . \quad (2.7)$$

Since the target organ T may receive dose from more than one source organ, the total value of H_{50} in target T is given by

$$H_{50,T} = 1.6 \times 10^{-10} \sum_s \sum_j \left[U_{s,j} \sum_i SEE(T \leftarrow S)_i \right]_j . \quad (2.8)$$

Since our interest is the specific committed dose equivalent, we express it per unit activity as

$$\hat{H}_{50,T} = 1.6 \times 10^{-10} \sum_s \sum_j \left[\hat{U}_{s,j} \sum_i SEE(T \leftarrow S)_i \right]_j . \quad (2.9)$$

where $\hat{U}_{s,j}$ is the number of transformations of radionuclide j in S over 50 years after intake of unit activity of the radionuclide.

2.1 SPECIFIC EFFECTIVE ENERGY

Let us denote the specific effective energy absorbed in target T per transformation of radionuclide j emitting all types of radiation i in source S as

$$SEE(T \leftarrow S)_j = \sum_i SEE(T \leftarrow S)_{i,j} . \quad (2.10)$$

As previously stated, each radionuclide may emit a range of different types and energies of radiation indicated by index i . But it is not necessary that every transformation (decay) of radionuclide j should result in emission of all the varied radiations. Hence the concept of yield. Associated with average or unique (as appropriate) energy E_i (in MeV) of every radiation type i is a yield Y_i . The yield provides the fraction of transformations that result in the release of radiation type i . Also, the amount of the energy absorbed in target T will vary significantly with the energy, kind of radiation, and location (source organ S) of release. This variance is accounted by introducing a quantity $AF(T \leftarrow S)_i$ which is defined as the fraction of energy absorbed in target organ T per emission of radiation i in source organ S . Of course, in accordance with the definition in the previous section, the product of all these quantities must be weighted for effective quality factors \bar{Q}_i for each radiation type i , and, since we are interested in specific effective energy, the whole product must be taken per unit mass M_T (in g) of the target organ T . We can thus write, for each radionuclide j ,

$$SEE(T \leftarrow S)_j = \sum_i \frac{Y_i E_i AF(T \leftarrow S)_i \bar{Q}_i}{M_T} . \quad (2.11)$$

- a) Decay Schemes [KOS1]: The decay schemes of radionuclides used in this report were obtained from "Radioactive Decay Data Tables," by D. C. Kocher, DOE/TIC - 11026(1981). In case of positron emission, a photon of energy 0.511 MeV and yield twice that of the

positron is added to the decay scheme to account for the annihilation photons. The decay scheme used in the ICRP-30 results were from Publication 38 [IC83].

- b) Absorbed fraction of energy in target organ: For most target organs, it is assumed that the energies from non-penetrating radiations such as alpha particles, beta particles, positrons, etc. are completely absorbed within the source organ. Hence, if target organ and source organ are the same, then the absorbed fraction is equal to 1, else it is equal to zero for non-penetrating radiation, except in the following cases [SN75]:

- i) If the source organ is total body then, regardless of the target organ, the specific absorbed fraction $\hat{A}F$ (absorbed fraction per g of target) is given by

$$\hat{A}F(\text{Total body}) = 1/69900,$$

where the denominator represents mass in grams of the total body of Reference Man.

- ii) When the target organ is total body and the source organ is either bladder content, stomach content, SI content, ULI content, or LLI content, then

$$\hat{A}F(\text{Total body} \leftarrow \text{GI tract/Bladder content}) = M_w / (2 \times M_c \times 69900),$$

where M_w is the mass of the wall of the source organs, and

M_c is the mass of the contents of the source organs.

If the source is any other organ excluding the above and the target organ is total body then

$$\hat{AF}(\text{Total body} \leftarrow S) = 1/69900.$$

With the specific absorbed fraction, the value of specific effective energy is given by

$$SEE(T \leftarrow S)_j = \sum_i Y_i E_i \hat{AF}(T \leftarrow S)_i \bar{Q}_i. \quad (2.12)$$

In case of penetrating radiations such as X-rays and γ rays, the absorbed fraction of energy from photons is estimated by the data in ICRP Publication 23 [IC75]. The tables in Publication 23 provide values for discrete energies from 0.01 MeV to 4.0 MeV. If the photon energy is within this range, then the value of absorbed fraction can be interpolated from the tables. However, if the energy of the photon is less than 0.01 MeV, specific absorbed fractions used in calculations are assumed to be the same as defined for non-penetrating radiations [SN78], i.e.,

- i) $\hat{AF} = 1/M_T$ if the source and the target are the same, or
- ii) $\hat{AF} = 0$ in general if the source and the target are different, except for the following conditions:
- iii) $\hat{AF} = 1/69900$ if the source organ is total body.
- iv) $\hat{AF} = M_w / (2 \times M_c \times 69900)$ if the target organ is total body and the source is either bladder content, stomach content, SI content, ULI content, or LLI content.
- v) $\hat{AF} = 1/69900$ if the target organ is total body and the source is any organ other than the ones described in (iv).

This treatment in calculation of absorbed fractions or specific absorbed fractions is recommended for the majority of target organs. However, there are exceptions, notably the mineral bone, organs of the GI tract, and the bladder wall, which are explained below:

2.1.1 Target Organs of the GI Tract and Bladder Wall:

For non-penetrating radiations, if the source organ is different from a target organ of the GI tract or bladder wall, the specific absorbed fraction is zero except when the source is total body. In that case, $\hat{A}F = 1/69900$. However, if the source is the content of the target organ in the GI tract or the bladder content, the only dose received is that by the mucosal layer ML of the wall of the organ. The specific absorbed fraction for the mucosal layer of the wall of the target organ is taken to be equal to $0.5 \times \nu / M_T^C$, where M_T^C (in g) is the mass of the contents of that target organ, and ν is a factor between zero and unity representing the degree to which the radiation penetrates the mucus. The factor 0.5 is introduced because the absorbed dose rate at the interface between the contents and the mucus and mucosal layers is approximately half that deep within the contents. For β particles, ν is taken to be unity, and for α particles only 0.01 owing to attenuation in the mucus layer.

As far as photons are concerned, the fraction of their energy emitted in source S that is absorbed in the walls of the target T is obtained from the tables in ICRP Publication 23 by interpolating within the 0.01 to 4.0 MeV range. However, if the energy of the photon is less than 0.01 MeV, the specific absorbed fraction is extrapolated as:

- i) $\hat{A}F = 1/(2 \times M_c)$ if the source and the target are the same, where M_c is the mass of contents of the organ, or
- ii) $\hat{A}F = 0$ in general if the source and the target are different, except in the following case:
- iii) if the source organ is total body, then $\hat{A}F = 1/69900$.

2.1.2 Target Organ in Bone:

In the case of mineral bone, the two target tissues are the cells near bone surfaces (BS) and the active red bone marrow (RM). The ICRP model describes source tissues as cortical and trabecular bone for all non-penetrating radiation. For photon emitters, S is any organ of the body containing the radionuclide and T is either the BS cells or RM. Cortical bone (CB) is the compact or dense material of the outside of the bone. Trabecular bone (TB) is the cancellous or spongy inner portion of the bone containing the marrow.

Absorbed fractions for non-penetrating radiation are governed by two major criteria:

- i) Radionuclides with half lives of less than 15 days are considered to be distributed on bone surfaces, since they are unlikely to move far into the volume of bone before they decay,
- ii) Isotopes of alkaline earth elements with half lives greater than 15 days and radionuclides ^{33}P , $^{93\text{m}}\text{Nb}$, ^{94}Nb , ^{232}U , ^{233}U , ^{234}U , ^{235}U , ^{236}U , ^{238}U , Na, Cr, Rb, ^{65}Zn , ^{205}Pb , ^{210}Pb , ^{49}V , ^7Be , ^{10}Be , ^{103}Pd , ^{107}Pd , ^{113}Sn , $^{119\text{m}}\text{Sn}$, ^{123}Sn , ^{126}Sn , ^{182}Ta ,

^{181}W , ^{185}W , ^{188}W are assumed to be uniformly distributed throughout the volume of bone.

If the source organ is any other organ except total body, trabecular bone, or cortical bone, the specific absorbed fraction is taken to be zero. If the source is total body, $\hat{AF} = 1/69900$. Recommended values of absorbed fractions for α and β particles in the cases of source tissues as TB and CB are given in Table 2.1. Masses of target organs BS and RM are taken to be 120 and 1500 g respectively.

TABLE 2.1 Absorbed fractions for dosimetry of radionuclides in bone as recommended by the ICRP 30.

Source Organ	Target Organ	α -emitter uniform in volume	α -emitter on bone surface	β -emitter uniform in volume	β -emitter on bone surface	β -emitter on bone surface
					$\bar{E} \geq 0.2 \text{ MeV}$	$\bar{E} < 0.2 \text{ MeV}$
TB	BS	0.025	0.25	0.025	0.025	0.25
CB	BS	0.01	0.25	0.015	0.015	0.25
TB	RM	0.05	0.5	0.35	0.5	0.5
CB	RM	0.0	0.0	0.0	0.0	0.0

For photon emitters, if the energy is within 0.01 to 4.0 MeV, values of absorbed fraction can be interpolated from the tables in ICRP Publication 23 for any source organ. Values of absorbed fraction reported for skeleton in Publication 23 are taken as appropriate for BS cells. If the photon energy is less than 0.01 MeV and the source is TB or CB, absorbed fraction is taken to be $1/M_T$, where M_T is 1500 g for

target organ RM and 10500 g for BS cells. If however, the source is any other organ except total body, $AF = 0$. In the case of total body, specific absorbed fraction $\hat{AF} = 1/69900$.

2.1.3 A note on daughters:

The metabolic behavior of daughters is assumed to be the same as that of the parent radionuclide which enters the body (see Section 2.2.1).

2.2 NUMBER OF TRANSFORMATIONS IN A SOURCE ORGAN OVER 50 YEARS

The number of transformations of a radionuclide in any source organ during a period of time is defined as the time integral of the activity of the radionuclide over that period of time.

After ingestion or inhalation of a radionuclide, its translocation to the body fluids is dependent upon the rate constants of the different compartments of the gastrointestinal and respiratory systems, and the decay constant of the radionuclide. A detailed account of the passage of radionuclides through the GI and respiratory system follows later. For now, let us examine the kinetics of a radionuclide after it reaches the body fluid compartment, and how it is deposited in, or passed for excretion from the different compartments of a tissue or organ. The mathematical model recommended by the ICRP is illustrated in Fig. 2.1. Transformations in the body fluid or transfer compartment are assumed to be uniformly distributed throughout the whole body of mass 70000 g. Each source organ or tissue may have one or more compartments where the radionuclide may be retained or translocated at

different rates. A maximum of three compartments per tissue is assumed. From each of these compartments, the radionuclide migrates at an appropriate rate to excretion pathways. In the interest of simplicity, no feedback to the transfer compartment either from the tissue compartments or from routes of excretion is assumed, although in reality the case is otherwise.

2.2.1 A note on daughters:

The immediate daughters and all subsequent progeny produced within the body are assumed to be associated with, and behave metabolically as the inhaled or ingested parent radionuclide. In general, there is little evidence to show whether the daughters behave metabolically like the parent or if, upon production, they exhibit their own metabolic behavior. When experimental evidence contrary to the assumption is available, separate models as shown later are used.

2.2.2 Transfer compartment, T:

Let us represent the radionuclides with an index j . If we suppose that an initial activity f_{T1}^j , per unit intake, reaches the body fluid or transfer compartment from the GI tract and lungs, then for the parent ($j = 1$) a differential equation expressing the time-dependent behavior in accordance with the model in Fig. 2.1, can be written as

$$\frac{dq_{T1}(t)}{dt} = -\lambda_T q_{T1}(t) - \lambda_1 q_{T1}(t) , \quad (2.13)$$

where q_{T1} is the activity of the parent in the transfer compartment,

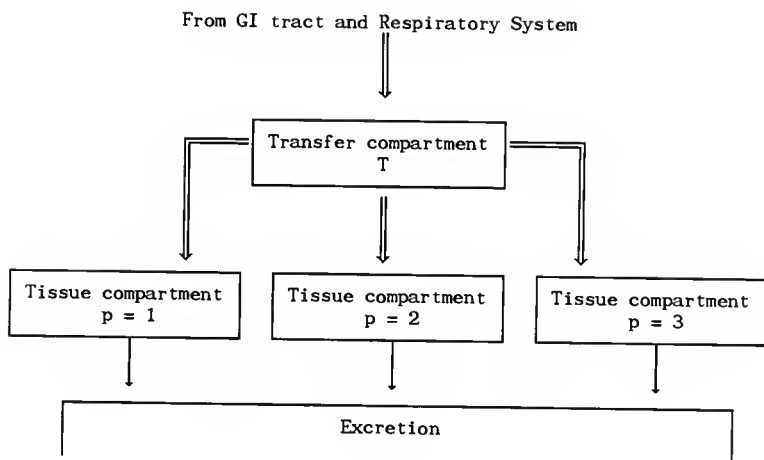


FIG 2.1 Mathematical model used by the ICRP-30 to describe the kinetics of radionuclides in the body.

λ_1 is the radiological decay constant for species 1, and

λ_T is the rate of loss of the stable element from the transfer compartment.

For most elements, λ_T is taken to be equal to $\ln 2 / 0.25$ days, i.e., the biological half life for translocation to the organs and tissues of deposition from the transfer compartment is taken to be 0.25 days. Following are the exceptions:

- i) If the element is fluorine, potassium, gold, or thallium, the translocation is assumed to be instantaneous and the biological half life is zero.

- ii) If the element is thorium, cobalt, chromium, or phosphorus, the biological half life is 0.5 days.
- iii) If the element is technetium or rhenium, the biological half life is 0.02 days.
- iv) If the element is ruthenium or rhodium, the biological half life is 0.3 days.
- v) If the element is tellurium, the biological half life is 0.8 days.
- vi) If the element is bismuth, the biological half life is 0.01 days.

Taking the Laplace transform of Eq. (2.13), we have

$$s \overline{q_{T1}}(s) - q_{T1}(0) = -\lambda_T \overline{q_{T1}}(s) - \lambda_1 \overline{q_{T1}}(s) .$$

This can be written as

$$\overline{q_{T1}}(s) = \frac{q_{T1}(0)}{[s + (\lambda_T + \lambda_1)]} . \quad (2.14)$$

Now for the daughters, a similar differential equation (see Fig. 2.1) can be written as

$$\frac{dq_{Tj}(t)}{dt} = \lambda_j q_{T,j-1}(t) - \lambda_T q_{Tj}(t) - \lambda_j q_{Tj}(t) ,$$

$$j = 2, \dots, N \quad (2.15)$$

where j is the index for the daughters,

q_{Tj} is the activity of the species j in the transfer compartment,

$q_{T,j-1}$ is the activity of the parent of species j in the transfer compartment, and

λ_j is the radiological decay constant for species j .

Specifically for $j = 2$,

$$\frac{dq_{T2}(t)}{dt} = \lambda_2 q_{T1}(t) - \lambda_T q_{T2}(t) - \lambda_2 q_{T2}(t) . \quad (2.16)$$

The Laplace transform of Eq. (2.16) is

$$s \overline{q_{T2}}(s) - q_{T2}(0) = \lambda_2 \overline{q_{T1}}(s) - \lambda_T \overline{q_{T2}}(s) - \lambda_2 \overline{q_{T2}}(s),$$

or

$$\overline{q_{T2}}(s) = \frac{q_{T2}(0) + \lambda_2 \overline{q_{T1}}(s)}{[s + (\lambda_T + \lambda_2)]} . \quad (2.17)$$

Substituting for $\overline{q_{T1}}(s)$ from Eq. (2.14) in Eq. (2.17), we obtain

$$\overline{q_{T2}}(s) = \frac{1}{[s + (\lambda_T + \lambda_2)]} \left\{ q_{T2}(0) + \frac{\lambda_2 q_{T1}(0)}{[s + (\lambda_T + \lambda_1)]} \right\} . \quad (2.18)$$

In general form,

$$\overline{q_{Tj}}(s) = \sum_{i=1}^j \frac{\left[\prod_{k=i+1}^j \lambda_k \right] q_{Ti}(0)}{\prod_{k=i}^j [s + (\lambda_T + \lambda_k)]} . \quad (2.19)$$

Now,

$$\frac{1}{\prod_{k=1}^j [s + (\lambda_T + \lambda_k)]} = \frac{1}{[s + (\lambda_T + \lambda_1)] [s + (\lambda_T + \lambda_2)] \dots}.$$

In partial fractions, this can be written as

$$\frac{1}{\prod_{k=1}^j [s + (\lambda_T + \lambda_k)]} = \frac{A_1}{s + (\lambda_T + \lambda_1)} + \frac{A_2}{s + (\lambda_T + \lambda_2)} + \dots + \frac{A_m}{s + (\lambda_T + \lambda_m)} + \dots$$

Multiplying throughout by $[s + (\lambda_T + \lambda_m)]$, we have

$$\frac{s + (\lambda_T + \lambda_m)}{\prod_{k=1}^j [s + (\lambda_T + \lambda_k)]} = \frac{A_m [s + (\lambda_T + \lambda_m)]}{[s + (\lambda_T + \lambda_m)]} + [s + (\lambda_T + \lambda_m)] \sum_{\substack{k=1 \\ k \neq m}}^j \frac{A_k}{[s + (\lambda_T + \lambda_k)]}.$$

Hence,

$$\frac{1}{\prod_{\substack{k=1 \\ k \neq m}}^j [s + (\lambda_T + \lambda_k)]} = A_m + [s + (\lambda_T + \lambda_m)] \sum_{\substack{k=1 \\ k \neq m}}^j \frac{A_k}{[s + (\lambda_T + \lambda_k)]}.$$

Let $s = -(\lambda_T + \lambda_m)$, then

$$A_m = \frac{1}{\prod_{\substack{k=1 \\ k \neq m}}^j (\lambda_k - \lambda_m)}.$$

Thus,

$$\overline{q_{Tj}}(s) = \sum_{i=1}^j \left\{ \left[\prod_{k=i+1}^j \lambda_k \right] q_{Ti}(0) \left[\sum_{\substack{m=i \\ k \neq m}}^j \frac{1}{\prod_{k=i}^j (\lambda_k - \lambda_m) [s + (\lambda_T + \lambda_m)]} \right] \right\}.$$

We know that if $L(f) = 1/s+a$, then $f(t) = e^{-at}$.

Therefore,

$$q_{Tj}(t) = \sum_{i=1}^j \left\{ \left[\prod_{k=i+1}^j \lambda_k \right] q_{Ti}(0) \left[\sum_{\substack{m=i \\ k \neq m}}^j \frac{e^{-(\lambda_T + \lambda_m)t}}{\prod_{k=i}^j (\lambda_k - \lambda_m)} \right] \right\}. \quad (2.20)$$

where $\prod_{i=m}^n a_i = a_m \times a_{m+1} \times \dots \times a_n$ if $n \geq m$, and

$$\prod_{i=m}^n a_i = 1 \text{ if } m > n.$$

Now by definition, the number of transformations in the transfer compartment U_{Tj} for species j is given by

$$U_{Tj} = B_j \times \int_0^T q_{Tj}(t) dt, \quad (2.21)$$

where $T = 365.25 \times 50$ days, and

B_j is the branching ratio of radionuclide j . For the parent, $B_1 = 1$.

The result of the integral is

$$U_{Tj} = B_j \times \sum_{i=1}^j \left\{ \left[\prod_{k=i+1}^j \lambda_k \right] q_{Ti}(0) \left[\sum_{m=i}^j \frac{[1 - e^{-(\lambda_T + \lambda_m)T}]}{\prod_{\substack{k=i \\ k \neq m}}^j (\lambda_T + \lambda_m)} \right] \right\}, \quad (2.22)$$

where $\prod_{i=m}^n a_i = a_m \times a_{m+1} \times \dots \times a_n$ if $n \geq m$.

$\prod_{i=m}^n a_i = 1$ if $m > n$, and

$q_{Ti}(0)$ is the initial activity of species i in the transfer compartment which is assumed to be equal to f_T^i .

Calculation of this quantity is shown later.

If λ values have units d^{-1} , then this formula for U_{Tj} must be multiplied by 86400 s/d to yield units Bq^{-1} for U_{Tj} .

2.2.3 Tissue compartment, p:

As mentioned earlier, each source organ or tissue can have a maximum of three compartments, i.e., $p = 1, 2, 3$. From the transfer compartment, a fraction f_2^p may reach each tissue compartment of a source organ. This fraction is biologically eliminated from the compartment with a half life of λ_p . Since the daughters are assumed to possess the same metabolic behavior as the parent, the same retention fractions f_2^p and biological half-lives λ_p are used for them. These retention fractions and biological half lives are obtained from the

metabolic data in ICRP Publication 30 for each nuclide taken in the body.

With the knowledge of f_2^P , the total activity of species j translocated from the transfer to the tissue compartment during the 50 year time period can be given by

$$Q_{pj} = f_2^P \lambda_T \int_0^T q_{Tj}(t) dt \quad . \quad (2.23)$$

But as shown before,

$$U_{Tj} = B_j \times \int_0^T q_{Tj}(t) dt \quad .$$

Hence,

$$Q_{pj} = \frac{f_2^P \lambda_T}{B_j} U_{Tj} \quad . \quad (2.24)$$

If we assume that this total activity is transferred instantaneously as a single intake to the tissue compartment at time $t = 0$, then the initial activity deposited in the tissue compartment $q_{pi}(0)$ for species i is equal to

$$q_{pi}(0) = \frac{f_2^P \lambda_T U_{Ti}}{B_i} \quad . \quad (2.25)$$

For the time dependent behavior of species j in the tissue compartment, differential equations similar to the ones described in

the transfer compartment can be written. By a similar treatment of solution, the number of transformations in a tissue compartment for species j can be written as

$$U_{pj} = B_j \times \sum_{i=1}^j \left\{ \left[\prod_{k=i+1}^j \lambda_k \right] q_{pi}(0) \left[\sum_{m=i}^j \frac{[1 - e^{-(\lambda_p + \lambda_m)T}]}{(\lambda_p + \lambda_m) \prod_{\substack{k=i \\ k \neq m}}^j (\lambda_k - \lambda_m)} \right] \right\} . \quad (2.26)$$

where $q_{pi}(0)$ is given by Eq. (2.25).

Thus, for a source which may have a maximum of 3 compartments, the number of source-organ transformations for species j is

$$U_{sj} = \frac{M_s \times U_{Tj}}{70000} + \sum_{p=1}^3 U_{pj} . \quad (2.27)$$

where M_s is mass of the source organ, and

70000 is mass of the total body in grams.

2.2.4 Respiratory system:

When a radionuclide is inhaled, parts of the respiratory system are irradiated. As a consequence, other organs and tissues of the body may be irradiated either by translocation of the inhaled material from the respiratory system to the body tissues or by radiations originating from the lungs.

Analyses of the dynamics of radionuclide transport within the lung is essential for the evaluation of number of source-organ transformations.

The model representing the respiratory system is proposed by the ICRP Task Group on Lung Dynamics (1966) [IC66]. It is shown in Fig. 2.2. This model partitions the respiratory system into three regions -- the nasal passage (N-P), the trachea and bronchial tree (T-B), and the pulmonary parenchyma (P). Each region is further subdivided into two or four compartments. All three regions have pathways directly to the body fluid compartment. Only the P region translocates material to the lymphatic system (L). This pulmonary lymphatic system also serves to remove the dust from the lungs. A subcompartment of the lymphatic system releases material to the body fluids while the other subcompartment is assumed to retain the material indefinitely. The latter subcompartment is deemed appropriate only for a particular class Y of aerosols. Both T-B and N-P regions are involved with mucociliary transport which translocates material to the gastrointestinal tract. Connection between the P region and the GI tract is only through feedback via the T-B region.

a) Deposition and Retention of Inhaled Material: Deposition of inhaled material in the respiratory system is dependent upon the aerodynamic properties of the aerosol distribution. Three terms D_{N-P} , D_{T-B} , and D_P represent the fractions of inhaled material initially deposited in the N-P, T-B, and P regions respectively, the balance being the fraction exhaled. The pattern of aerosol distribution is characterized by the activity median aerodynamic diameter (AMAD). This quantity is closely approximated by the mass-median aerodynamic

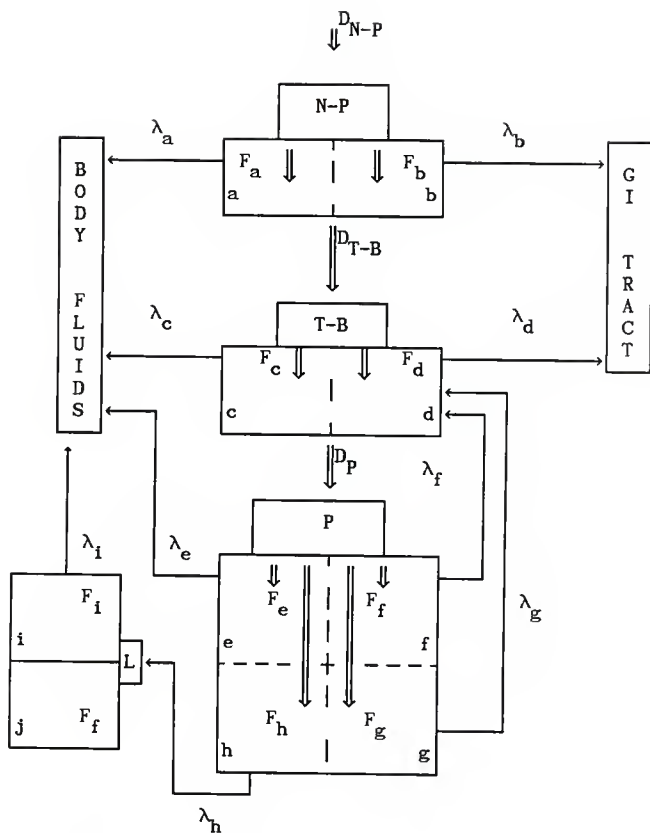


FIG. 2.2. Mathematical model used to describe clearance from respiratory system.

Source: ICRP Publication 30, Part I.

diameter, the aerodynamic diameter of a particle being the diameter of a spherical particle with the same settling velocity. The pattern of deposition is related to the AMAD of the aerosol according to Fig. 2.3.

To analyze the clearance of inhaled radioactive materials from the lung, the radionuclides are categorized as D, W, or Y. These categories refer to the retention in the pulmonary region. Class D reflects minimal retention, i.e., a rapid clearance within days while, class W material takes weeks for clearance, and class Y is cleared slowly in years. For elements and their compounds, retention classification is given in Fig. 2.4.

For each chemical classification, the sub-compartments have a half time of clearance T (days), and a fraction F that refers to the amount of material leaving each sub-compartment. These values are shown in Table 2.2. Note that the half lives are given in units of days. When converting these decay constants in units of s^{-1} , one must use the factor 86400 s/d.

b) Clearance Calculations: To calculate the source-organ transformations of a nuclide, we need to evaluate its time dependent activity in each subcompartment. Suppose we represent the subcompartments with an index ℓ such that $\ell = a, b, c, d, e, f, g, h, i$, and j as shown in Fig. 2.2. Now, if a unit activity of nuclide is inhaled at time $t = 0$, then the initial activities in each of the subcompartments $a - h$ of Fig. 2.2 can be given as the product of percent deposition in the compartment and the fraction of material entering the

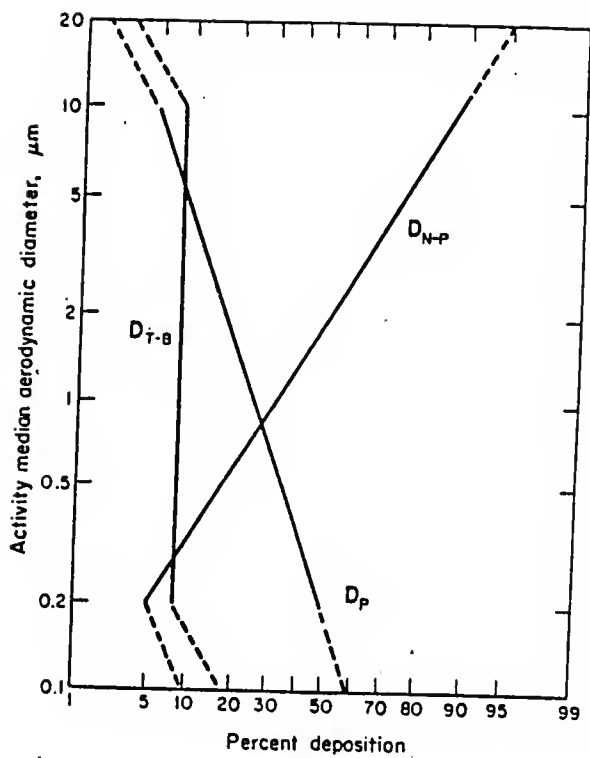


FIG. 2.3. Model of deposition of dust in the respiratory system as proposed by the ICRP-30. The model is intended for use with aerosols with AMAD between 0.2 and 10 μm and with geometric standard deviations of less than 4.5. Provisional estimates of deposition further extending the size range and given by dashed lines.

Class F—Arid retention: cleared slowly (years)
Carbides—actinides, lanthanides, Zr, Y, Mn
Sulfides—none
Sulfates—none
Carbonates—none
Phosphates—none
Oxides and hydrides—lanthanides, actinides Groups 8 (V and VI), 1b, 2b (IV and V), 3b except Sc ⁴⁺ and 6b.
Halides—lanthanide fluorides
Nitrides—none
Class W—Moderate retention: intermediate clearance rates (weeks)
Carbides—Cations of all Class W hydrides except those listed as Class Y carbides.
Sulfides—Groups 2a (V + VI), 4a (IV-VI), 5a (IV-VI), 1b, 2b and 6b (V + VI).
Sulfates—Groups 2a (IV-VI), and 3a (IV-VI)
Carbonates—lanthanides, Bi ³⁺ and Group 2a (IV-VI)
Phosphates—Zn ²⁺ , Sn ²⁺ , Mg ²⁺ , Fe ²⁺ , Bi ³⁺ and lanthanides
Oxides and hydrides—Groups 2a (II-VII), 3a (III-VI), 4a (III-VI), 5a (IV-VI), 6a (IV-VI), 8, 2b (VI), 4b, 5b, and 7b Sc ³⁺
Halides—lanthanides (except fluorides), Groups 2a, 3a (III-VI), 4a (IV-VI), 5a (IV-VI), 6a (IV-VI), 8, 2b (IV-V), 4b, 5b, 6b and 7b
Nitrides—all cations whose hydrides are Class Y and W
Class D—Minimal retention: rapid clearance (days)
Carbides—see hydrides
Sulfides—all except Class W
Sulfates—all except Class W
Carbonates—all except Class W
Phosphates—all except Class W
Oxides and Hydrides—Groups 1a, 3a (II), 4a (II), 5a (II, III), 6a (III).
Halides—Groups 1a and 7a
Nitrides—all except Class W
Noble Gases—Group 0

Note: Where reference is made from one chemical form to another, it implies that an *in vivo* conversion occurs, e.g. hydrolysis reaction.

The following periodic table of the elements is used with the foregoing classification.

Period	Group															
	1a	2a	3b	4b	5b	6b	7b	8	9	10	11	12	13	14	15	16
I	H															He
II	Li	Be											B	C	N	O
III	Na	Mg											Al	Si	P	S
IV	K	Ca	Sc	Ti	V	Cr	Mn	Fe	Co	Ni	Cu	Zn	Ga	Ge	As	Se
V	Rb	Sr	Y	Zr	Nb	Mo	Tc	Ru	Rh	Pd	Ag	Cd	In	Sn	Sb	Te
VI	Cs	Ba	La*	Hf	Ta	W	Re	Os	Ir	Pt	Au	Hg	Tl	Pb	Bi	Po
VII	Fr	Ra	Ac†													
* Lanthanides																
	Ca	Fr	Nd	Pm	Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb	Lu		
† Actinides																
	Th	Pa	U	Np	Pu	Am	Cm	Bk	Cf	Es	Fm	Md	No	Lw		

FIG. 2.4. Pulmonary clearance classification of inorganic compounds as proposed by the ICRP.

Source: Report of ICRP Task Group on Lung Dynamics: Health Physics, 12, 173-207 (1966).

TABLE 2.2. The values of removal half times and compartmental fractions in the four respiratory regions as proposed in ICRP-30.

		Class					
		D		W		Y	
		T day	F	T day	F	T day	F
N-P ($D_{N-P} = 0.30$)	a	0.01	0.5	0.01	0.1	0.01	0.01
	b	0.01	0.5	0.40	0.9	0.40	0.99
T-B ($D_{T-B} = 0.08$)	c	0.01	0.95	0.01	0.5	0.01	0.01
	d	0.2	0.05	0.2	0.5	0.2	0.99
P ($D_P = 0.25$)	e	0.5	0.8	50	0.15	500	0.05
	f	n.a.*	n.a.	1.0	0.4	1.0	0.4
	g	n.a.	n.a.	50	0.4	500	0.4
	h	0.5	0.2	50	0.05	500	0.15
L	i	0.5	1.0	50	1.0	1000	0.9
	j	n.a.	n.a.	n.a.	n.a.	∞	0.1

* n.a. = not applicable.

subcompartment. For example, the initial activity in subcompartment d is $q_d(0) = D_{T-B} F_d$. The clearance of the material from each of the subcompartments is assumed to be governed by first order kinetics, so that each subcompartment is associated with a biological decay constant, e.g., $\lambda_d(s^{-1}) = \ln 2 / (86400 T_d)$. Activities in each subcompartment may be found by solving the first order differential equations as follows:

For $\ell = a, b, c, e, f, g,$ and h ,

$$\frac{dq_{\ell}}{dt} = - (\lambda_R + \lambda_{\ell}) q_{\ell} . \quad (2.28)$$

Else,

$$\frac{dq_d}{dt} = \lambda_f q_f + \lambda_g q_g - (\lambda_R + \lambda_d) q_d , \quad (2.29)$$

$$\frac{dq_i}{dt} = - (\lambda_R + \lambda_i) q_i + F_i \lambda_h q_h , \quad (2.30)$$

$$\frac{dq_j}{dt} = - \lambda_R q_j + F_j \lambda_h q_h . \quad (2.31)$$

Since daughters are assumed to behave metabolically like the inhaled parent, similar systems of equations may be written for them using the partition fractions F and biological half lives as those of the parent.

The approximate expressions for the total number of transformations of the parent and the daughters in each subcompartment of the lung is given in Tables 2.3 and 2.4 respectively. These results are obtained by integrating the activities found from the solution of the above equations, in a pattern similar to that of the transfer compartment. All entries, except that for subcompartment j , are based on the approximation that the clearance times are much less than the 50-year dose evaluation period. In the entry for compartment j , the symbol η represents 50 years in units compatible with those of λ_R .

TABLE 2.3. Approximate expressions for the number of transformations in the subcompartments of lung following inhalation of unit Bq of activity. (a)

Compartment	Number of transformations	Compartment	Number of transformations
a	$\frac{D_{N-P} F_a}{\lambda_a + \lambda_R}$	b	$\frac{D_{N-P} F_b}{\lambda_b + \lambda_R}$
c	$\frac{D_{T-B} F_c}{\lambda_c + \lambda_R}$	d	$\left[\frac{D_{T-B} F_d}{\lambda_d + \lambda_R} + \frac{D_p}{\lambda_d + \lambda_R} \left[\frac{\lambda_f F_f}{\lambda_f + \lambda_R} + \frac{\lambda_g F_g}{\lambda_g + \lambda_R} \right] \right]$
e	$\frac{D_p F_e}{\lambda_e + \lambda_R}$	f	$\frac{D_p F_f}{\lambda_f + \lambda_R}$
g	$\frac{D_p F_g}{\lambda_g + \lambda_R}$	h	$\frac{D_p F_h}{\lambda_h + \lambda_R}$
i	$\frac{D_p F_h \lambda_h F_i}{(\lambda_h + \lambda_R)(\lambda_i + \lambda_R)}$	j	$\frac{D_p F_h \lambda_h F_j (1 - e^{-\eta \lambda_R})}{(\lambda_h + \lambda_R) \lambda_R}$

(a) Source: ICRP Publication 30 [IC79].

TABLE 2.4. Approximate expressions for the number of transformations of a radioactive daughter in the subcompartments of the lung. A_a to A_j refers to the number of transformations of the immediate parent in subcompartments a to j of the lung. λ'_R is the radiological decay constant of the daughter. (a)

Compartment	Number of transformations	Compartment	Number of transformations
a	$\frac{A_a \lambda'_R}{\lambda_a + \lambda'_R}$	b	$\frac{A_b \lambda'_R}{\lambda_b + \lambda'_R}$
c	$\frac{A_c \lambda'_R}{\lambda_c + \lambda'_R}$	d	$\frac{A_d \lambda'_R}{\lambda_d + \lambda'_R} + \frac{\lambda'_R}{\lambda_d + \lambda'_R} \left[\frac{A_f \lambda'_R}{\lambda_f + \lambda'_R} + \frac{A_g \lambda'_R}{\lambda_g + \lambda'_R} \right]$
e	$\frac{A_e \lambda'_R}{\lambda_e + \lambda'_R}$	f	$\frac{A_f \lambda'_R}{\lambda_f + \lambda'_R}$
g	$\frac{A_g \lambda'_R}{\lambda_g + \lambda'_R}$	h	$\frac{A_h \lambda'_R}{\lambda_h + \lambda'_R}$
i	$\frac{A_i \lambda'_R}{\lambda_i + \lambda'_R} + \frac{A_h \lambda'_R \lambda_h F_i}{(\lambda_h + \lambda'_R)(\lambda_i + \lambda'_R)}$	j	$\left[A_j + \frac{A_h \lambda_h F_j}{\lambda_h + \lambda'_R} \right] \left[1 - e^{-\eta \lambda'_R} \right]$

(a) Source: ICRP Publication 30 [IC79].

With the source organ transformations known in each subcompartment, we can sum them all to determine a total value for the respiratory system. However, the ICRP model neglects the values for transformations in the nasopharyngeal region, since for most particle sizes, the dose received by this region is very small compared to the other regions. Hence, defining the lung as a single source organ, the value of transformations for a nuclide k is given by

$$U_L^k = [U_{T-B}^k + U_P^k + U_L^k] \times B_k, \quad (2.32)$$

where $U_{T-B}^k = U_C^k + U_D^k,$

$$U_P^k = U_e^k + U_f^k + U_g^k + U_h^k,$$

$$U_L^k = U_1^k + U_j^k, \text{ and}$$

B_k is the branching ratio of nuclide k ($B_1 = 1$).

The lung is also treated as a single target organ of mass 1000 g.

c) Transfer of a Radionuclide from the Lungs Directly to Body Fluids or to the GI Tract: From Fig. 2.2 we can see that the rate of transfer of a radionuclide k directly from the lungs to the body fluids [BF(t)] is given by

$$BF^k(t) = \lambda_a q_a^k(t) + \lambda_c q_c^k(t) + \lambda_e q_e^k(t) + \lambda_i q_i^k(t), \quad (2.33)$$

whether the radionuclide is inhaled or produced in lungs. Similarly, the rate of transfer to the GI tract is given by

$$G^k(t) = \lambda_b q_b^k(t) + \lambda_d q_d^k(t) . \quad (2.34)$$

Thus, the total activity f_{BFDIR}^k of an inhaled radionuclide k transferred directly to the body fluids can be determined as

$$f_{\text{BFDIR}}^k = B_k \times \int_0^{50y} Bf^k(t) dt , \quad (2.35)$$

or

$$f_{\text{BFDIR}}^k = B_k \times \left[\lambda_a \int_0^{50y} q_a^k(t) dt + \lambda_c \int_0^{50y} q_c^k(t) dt + \lambda_e \int_0^{50y} q_e^k(t) dt + \lambda_i \int_0^{50y} q_i^k(t) dt \right] , \quad (2.36)$$

or

$$f_{\text{BFDIR}}^k = B_k \times \left[\lambda_a U_a^k + \lambda_c U_c^k + \lambda_e U_e^k + \lambda_i U_i^k \right] . \quad (2.37)$$

Similarly, the total activity translocated to the gastrointestinal tract is

$$f_{\text{GI}}^k = B_k \times \left[\lambda_b U_b^k + \lambda_d U_d^k \right] . \quad (2.38)$$

d) Particle Size Correction: If the AMAD is unknown, then a value of $1 \mu\text{m}$ is used for inhaled materials. Values of specific committed dose are reported for $1 \mu\text{m}$ in ICRP Publication 30. Correction for other

values of AMAD is made as follows:

$$\frac{\hat{H}_{50,T}(\text{AMAD})}{\hat{H}_{50,T}(1\mu\text{m})} = f_{N-P} \frac{D_{N-P}(\text{AMAD})}{D_{N-P}(1\mu\text{m})} + f_{T-B} \frac{D_{T-B}(\text{AMAD})}{D_{T-B}(1\mu\text{m})} + f_P \frac{D_P(\text{AMAD})}{D_P(1\mu\text{m})}$$

The fractions f_{N-P} , f_{T-B} , and f_P are respectively the proportions of $\hat{H}_{50,T}(1\mu\text{m})$ resulting from deposition in the N-P, T-B, and P regions.

If the AMAD is not $1\mu\text{m}$, then the values of $D_{N-P}(\text{AMAD})$, $D_{T-B}(\text{AMAD})$, and $D_P(\text{AMAD})$ are determined from a logarithm fit of Fig. 2.3 as described below:

If the AMAD is between 0.1 and $0.2\mu\text{m}$, then

$$D_{T-B} = -0.163 - 0.151 (\ln \text{AMAD}), \quad (2.39)$$

$$D_{N-P} = -0.059 - 0.068 (\ln \text{AMAD}), \text{ and} \quad (2.40)$$

$$D_P = 0.289 - 0.126 (\ln \text{AMAD}) . \quad (2.41)$$

If the AMAD is between 0.2 and $10\mu\text{m}$, then

$$D_{T-B} = 0.08 , \quad (2.42)$$

$$D_{N-P} = 0.351 + 0.219 (\ln \text{AMAD}), \text{ and} \quad (2.43)$$

$$D_P = 0.289 - 0.126 (\ln \text{AMAD}) . \quad (2.44)$$

If the AMAD is between 10 and 20 μm , then

$$D_{T-B} = 0.229 - 0.065 (\ln \text{AMAD}) , \quad (2.45)$$

$$D_{N-P} = 0.621 + 0.110 (\ln \text{AMAD}), \text{ and} \quad (2.46)$$

$$D_P = 0.141 - 0.040 (\ln \text{AMAD}) . \quad (2.47)$$

2.2.5 Gastrointestinal Tract:

Knowledge of transformations of a nuclide in the various organs of the GI tract is very crucial to the calculation of other source-organ transformations, because translocation of the material to the body fluid compartment and then eventually to the source organ is directly dependent, in case of ingestion, and partially dependent, in case of inhalation, upon absorption of the material in the bloodstream through the small intestine.

The model proposed by the ICRP is illustrated in Fig. 2.5. It consists of 4 organs. Table 2.5 provides values of wall mass, contents mass, and mean residence times for the contents of each organ. The rate constant λ for transfer of contents from organ to organ is the reciprocal of the mean residence time.

In general, the only site of absorption from the GI tract to the body fluids is assumed to be the small intestine. The rate constant λ_{BF} for transfer from the small intestine to the body fluids after ingestion, can be estimated from the fraction of stable element

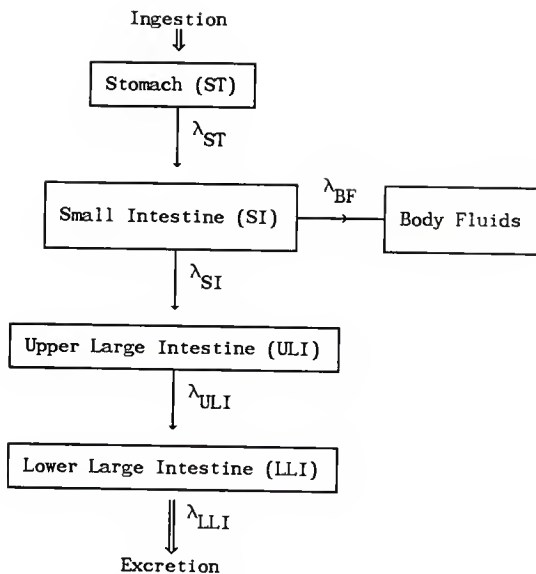


FIG. 2.5. Model of gastrointestinal tract as proposed by the ICRP-30.

reaching the body fluids f_1 , i.e.,

$$\frac{\lambda_{BF}}{\lambda_{SI} + \lambda_{BF}} = f_1 \quad (2.48)$$

Therefore,

$$\lambda_{BF} = \frac{f_1 \lambda_{SI}}{1 - f_1} \quad (2.49)$$

Values of f_1 for a number of classes of compounds of each element are given in the metabolic data. Radioactive progeny are assumed to have the same f_1 as the ingested ancestral nuclide.

When a value of $f_1 = 1$ is given, it implies that the radionuclide passes directly from the stomach to body fluids and does not pass through other sections of the GI tract.

TABLE 2.5. Mathematical model used to describe the kinetics of radionuclides in the gastrointestinal tract as proposed by the ICRP-30.

Section of Tract	Mass of Walls (g)	Mass of Contents (g)	Mean Residence time (day)	λ (day ⁻¹)
Stomach (ST)	150	250	1/24	24
Small Intestine (SI)	640	400	4/24	6
Upper Large Intestine (ULI)	210	220	13/24	1.8
Lower Large Intestine (LLI)	160	135	24/24	1

a) Radionuclide Transport in the GI Tract: Consider the ingestion of unit activity of a radionuclide at time $t = 0$. For this parent

radionuclide, the time dependent activity in different sections of the GI tract is governed by the following equations:

$$\frac{dq_{ST}(t)}{dt} = - (\lambda_{ST} + \lambda_R) q_{ST}(t) , \quad (2.50)$$

$$\frac{dq_{SI}(t)}{dt} = - (\lambda_R + \lambda_{SI} + \lambda_{BF})q_{SI}(t) + \lambda_{ST} q_{ST}(t) , \quad (2.51)$$

$$\frac{dq_{ULI}(t)}{dt} = - (\lambda_{ULI} + \lambda_R) q_{ULI}(t) + \lambda_{SI} q_{SI}(t) , \quad (2.52)$$

$$\frac{dq_{LLI}(t)}{dt} = - (\lambda_R + \lambda_{LLI}) q_{LLI}(t) + \lambda_{ULI} q_{ULI}(t) , \quad (2.53)$$

where λ_R is the radiological decay constant of the radionuclide, and

$\lambda_{BF}q_{SI}(t)$ is the rate of transfer of activity to the body fluids from the small intestine.

A similar set of equations may be written for the radioactive progeny. Solution of these equations can then be used in calculating the number of transformations in each section of the GI tract. Approximate expressions for numbers of transformations of the parent and daughters in the various sections of the tract, following ingestion of unit activity, is given in Tables 2.6 and 2.7 respectively. The approximation is that the residence times are short in comparison to the 50 year dose-evaluation time. These values should, of course, be multiplied by the corresponding branching ratio of the radionuclide.

TABLE 2.6. Approximate expressions for the number of transformations of the parent in the various regions of the gastrointestinal tract following ingestion of 1 Bq of activity. Based on ICRP Publication 30.

Region	Number of Transformations
Stomach	$\frac{1}{\lambda_{ST} + \lambda_R}$
Small Intestine	$\frac{\lambda_{ST}}{(\lambda_{ST} + \lambda_R)(\lambda_{SI} + \lambda_{BF} + \lambda_R)}$
Upper Large Intestine	$\frac{\lambda_{ST} \lambda_{SI}}{(\lambda_{ST} + \lambda_R)(\lambda_{SI} + \lambda_{BF} + \lambda_R)(\lambda_{ULI} + \lambda_R)}$
Lower Large Intestine	$\frac{\lambda_{ST} \lambda_{SI} \lambda_{ULI}}{(\lambda_{ST} + \lambda_R)(\lambda_{SI} + \lambda_{BF} + \lambda_R)(\lambda_{ULI} + \lambda_R)(\lambda_{LLI} + \lambda_R)}$

TABLE 2.7. Approximate expressions for the number of transformations of a radioactive daughter in the various sections of the GI tract. A_{ST} , A_{SI} , A_{ULI} , A_{LLI} are the number of transformations of the parent in the various regions of the tract. λ'_R is the radiological decay constant of the daughter. Based on ICRP Publication 30.

Region	Number of Transformations
Stomach	$\frac{A_{ST} \lambda'_R}{\lambda_{ST} + \lambda'_R}$
Small Intestine	$\frac{A_{ST} \lambda'_R \lambda_{ST}}{(\lambda_{ST} + \lambda'_R)(\lambda_{SI} + \lambda_{BF} + \lambda'_R)} + \frac{A_{SI} \lambda'_R}{(\lambda_{SI} + \lambda_{BF} + \lambda'_R)}$
Upper Large Intestine	$\frac{A_{ST} \lambda'_R \lambda_{ST} \lambda_{SI}}{(\lambda_{ST} + \lambda'_R)(\lambda_{SI} + \lambda_{BF} + \lambda'_R)(\lambda_{ULI} + \lambda'_R)}$ $+ \frac{A_{SI} \lambda'_R \lambda_{SI}}{(\lambda_{SI} + \lambda_{BF} + \lambda'_R)(\lambda_{ULI} + \lambda'_R)} + \frac{A_{ULI} \lambda'_R}{(\lambda_{ULI} + \lambda'_R)}$
Lower Large Intestine	$\frac{A_{ST} \lambda'_R \lambda_{ST} \lambda_{SI} \lambda_{ULI}}{(\lambda_{ST} + \lambda'_R)(\lambda_{SI} + \lambda_{BF} + \lambda'_R)(\lambda_{ULI} + \lambda'_R)(\lambda_{LLI} + \lambda'_R)}$ $+ \frac{A_{SI} \lambda'_R \lambda_{SI} \lambda_{ULI}}{(\lambda_{SI} + \lambda_{BF} + \lambda'_R)(\lambda_{ULI} + \lambda'_R)(\lambda_{LLI} + \lambda'_R)}$ $+ \frac{A_{ULI} \lambda'_R \lambda_{ULI}}{(\lambda_{ULI} + \lambda'_R)(\lambda_{LLI} + \lambda'_R)} + \frac{A_{LLI} \lambda'_R}{(\lambda_{LLI} + \lambda'_R)}$

b) Inhaled Radionuclides: In case of inhaled radionuclides, the radioactive material is translocated from the various subcompartments of the lung to the GI tract. The total activity transferred is different for different nuclides and is given by f_{GI}^k for species k . In case of the parent, the number of transformations in the various sections of the GI tract can simply be found by multiplying the expressions in Table 2.6 by f_{GI}^1 ($k = 1$ for parent). However, in case of the radioactive daughters, the expressions in Table 2.7 cannot be used, since each daughter may be deposited initially in the GI tract in different amounts, i.e., f_{GI}^k is different, and the immediate parent may decay into the daughter in the sections of the tract.

For this different set of differential equations for activity in the GI tract, compared to the ones in the case of ingestion, the number of transformations, derived in the pattern described in the transfer compartment, results in

$$U_{Gj} = B_j \times \sum_{i=1}^j \left\{ \left[\prod_{k=i+1}^j \lambda_k \right] q_{Gi}(0) \left[\sum_{m=1}^j \frac{(1 - e^{-(\lambda_G + \lambda_m)T})}{(\lambda_G + \lambda_m) \prod_{\substack{k=i \\ k \neq m}}^j (\lambda_k - \lambda_m)} \right] \right\}, \quad (2.54)$$

$$j = 2, \dots, N$$

where

G is the index for the different sections of the tract, i.e., ST, SI, ULI, and LLI.

T is, of course, the 50 year period.

λ_G is the clearance constant corresponding to the section of the GI tract. For example, in case of the stomach, $\lambda_G = \lambda_{ST}$. However, in case of small intestine,

$$\lambda_G = \lambda_{SI} + \lambda_{BF} .$$

j is in the index of the radioactive progeny of the parent, and

$q_{Gi}(0)$ is the total activity deposited instantaneously at time $t = 0$.

The last term is given by the following expression

$$q_{Gi}(0) = \lambda_{G-1,i} \int_0^{50y} q_{G-1,i}(t) dt, \quad (2.55)$$

where $G-1$ represents the compartment immediately preceding the one for which the transformations are calculated.

But,

$$U_{G-1,i} = \left[\int_0^{50y} q_{G-1,i}(t) dt \right] \times B_i, \quad (2.56)$$

Therefore,

$$q_{Gi}(0) = \lambda_{G-1,i} U_{G-1,i} / B_i. \quad (2.57)$$

$U_{G-1,i}$ is the number of transformations of species i in the preceding compartment. For example, in the case of calculation of the number of transformations in the small intestine, $U_{G-1,i} = U_{ST,i}$ and in the case of stomach, $U_{G-1,i} = f_{GI}^i$.

2.2.6 Activity translocated to the transfer compartment f_T^i :

This quantity was assumed to be known in the calculation of number of transformations in the transfer compartment. After the discussion

of the respiratory and the gastrointestinal system, we are now in a position to calculate this quantity.

a) Ingestion: In the case of ingestion of radionuclides, the total activity of species i translocated to the transfer compartment from the small intestine, per unit activity ingested, i is given by,

$$f_T^i = \int_0^{50y} \lambda_{BF} q_{SI}^i(t) dt, \quad (2.58)$$

or

$$f_T^i = \lambda_{BF} U_{SI}^i, \quad (2.59)$$

where U_{SI}^i is given in Tables 2.6 and 2.7 for the parent and the daughters respectively.

If the fraction of the stable element reaching the body fluids f_1 is 1, then the translocation is considered to be directly from the stomach. In this case,

$$f_T^i = \lambda_{ST} U_{ST}^i. \quad (2.60)$$

b) Inhalation: In case of inhalation, there are two pathways to the body fluid compartment. One is directly from the different compartments of the respiratory system and the other is indirectly through the material deposited in the GI tract from the respiratory system. Hence, the total initial activity f_T^i of species i deposited in the transfer compartment is given by

$$f_T^i = \left\{ f_{\text{BFDIR}}^i + f_{\text{GI}}^i \left[\lambda_{\text{BF}} U_{\text{SI}}^i \right] \right\} \times \frac{1}{B_i}, \quad (2.61)$$

where B_i is the branching ratio of species i ,

f_{BFDIR}^i is the total activity of inhaled radionuclide i transferred directly to the body fluid compartment, which is given by equation (2.37), and

f_{GI}^i is the total activity translocated to the gastrointestinal tract, given by equation (2.38).

Again, in case of $f_1 = 1$, the above equation can be modified as

$$f_T^i = \left\{ f_{\text{BFDIR}}^i + f_{\text{GI}}^i \left[\lambda_{\text{ST}} U_{\text{ST}}^i \right] \right\} \times \frac{1}{B_i} \quad (2.62)$$

2.2.7 Source organ as bone:

According to the ICRP model, the transformations in the cortical and trabecular bone are taken as a fraction of the transformations in the mineral bone calculated by the methods described earlier for any general source organ.

- i) For the parent radionuclide assumed to be on bone surface,
 $U_{\text{trabecular}}^i = U_{\text{cortical}}^i = 0.5 U_{\text{mineral bone}}^i$ for any species i including the radioactive progeny.
- ii) For the parent radionuclide assumed to be uniformly distributed throughout the volume of mineral bone,
 $U_{\text{trabecular}}^i = 0.2 U_{\text{mineral bone}}^i$ and $U_{\text{cortical}}^i = 0.8 U_{\text{mineral bone}}^i$ for any species i .

2.2.8 Three compartment model for iodine:

Iodine and all its radioactive isotopes require an exception to the treatment described for all other radionuclides. The metabolic model proposed for iodine by the ICRP is shown in Fig. 2.6. The model consists of three compartments. Since the value of $f_1 = 1$ for isotopes of iodine, the material is translocated from the stomach to the body fluids.

Of iodine entering the body fluid compartment, a fraction 0.3 is assumed to be translocated to the thyroid while the remainder is assumed to go directly to excretion. Iodine in the thyroid is assumed to be retained with a biological half-life of 120 days and to be lost

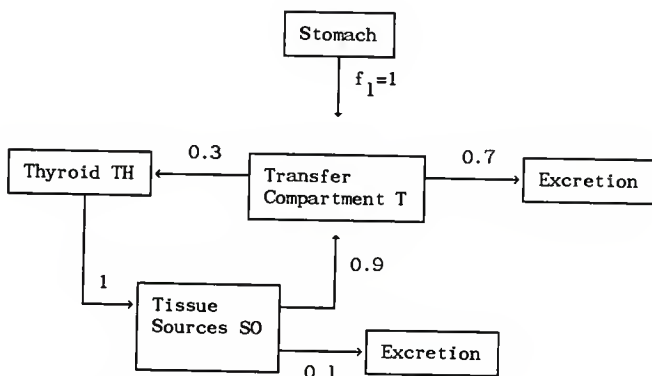


FIG. 2.6. Mathematical model proposed by the ICRP-30 for radioactive iodine.

from the gland in the form of organic iodine. Organic iodine is assumed to be uniformly distributed among all organs and tissues of the body other than the thyroid and to be retained there with a biological half-life of 12 days. One-tenth of this organic iodine is assumed to go directly to faecal excretion and the rest is assumed to be returned to the transfer compartment as inorganic iodine.

The time-dependent activity of the nuclide in the three compartments in accordance with Fig. 2.6 is given by the following differential equation for species i:

$$\frac{dq_{T,i}(t)}{dt} = -\lambda_T q_{T,i}(t) - \lambda_i q_{T,i}(t) + 0.9 \lambda_{SO} q_{SO,i}(t), \quad (2.63)$$

$$@ t = 0, \quad q_{T,i}(0) = f_T^i,$$

$$\lambda_{SO} = \ln 2/12 \text{ days}.$$

$$\frac{dq_{Th,i}(t)}{dt} = 0.3 \lambda_T q_{T,i}(t) - \lambda_{Th} q_{Th,i}(t) - \lambda_i q_{Th,i}(t) \quad (2.64)$$

$$@ t = 0, \quad q_{Th,i}(0) = 0,$$

$$\lambda_{Th} = \ln 2/120 \text{ days}.$$

$$\frac{dq_{SO,i}(t)}{dt} = \lambda_{Th} q_{Th,i}(t) - \lambda_{SO} q_{SO,i}(t) - \lambda_i q_{SO,i}(t), \quad (2.65)$$

$$@ t = 0, \quad q_{SO,i}(0) = 0.$$

Indices S0, T, and TH represent tissue sources, body fluid, and thyroid respectively. Taking the Laplace transform of the above equations, we obtain

$$\overline{q_{T,i}}(s) = \frac{0.9 \lambda_{S0} \overline{q_{S0,i}}(s) + f_T^i}{[s + (\lambda_T + \lambda_i)]} , \quad (2.66)$$

$$\overline{q_{Th,i}}(s) = \frac{0.3 \lambda_T \overline{q_{T,i}}(s)}{[s + (\lambda_i + \lambda_{Th})]} , \quad (2.67)$$

$$\overline{q_{S0,i}}(s) = \frac{\lambda_{Th} \overline{q_{S0,i}}(s)}{[s + (\lambda_i + \lambda_{S0})]} . \quad (2.68)$$

Since we are interested in the number of transformations in each of the compartments, which is the 50 year integral of the activities, we can use the final value theorem as follows:

$$\text{Let } F(t) = \int_0^t q(t') dt' , \text{ then } U = \lim_{t \rightarrow \infty} F(t) = \lim_{s \rightarrow 0} s \overline{F}(s) .$$

But

$$\overline{F}(s) = \frac{1}{s} \overline{q}(s) ,$$

therefore,

$$U = \lim_{s \rightarrow 0} \overline{q}(s) .$$

Using Cramer's rule, we obtain

$$U_T^i = \frac{f_T^i (\lambda_i + \lambda_{SO})(\lambda_i + \lambda_{Th})}{(\lambda_i + \lambda_T)(\lambda_i + \lambda_{Th})(\lambda_i + \lambda_{SO}) - 0.3(0.9)\lambda_T \lambda_{SO} \lambda_{Th}} . \quad (2.69)$$

$$U_{Th}^i = \frac{0.3 \lambda_T f_T^i (\lambda_i + \lambda_{SO})}{(\lambda_i + \lambda_T)(\lambda_i + \lambda_{Th})(\lambda_i + \lambda_{SO}) - 0.3(0.9)\lambda_T \lambda_{SO} \lambda_{Th}} + \frac{M_{Th} \times U_T^i}{70000}, \quad (2.70)$$

$$U_{SO}^i = \frac{0.3 \lambda_T \lambda_{Th} f_T^i}{(\lambda_i + \lambda_T)(\lambda_i + \lambda_{Th})(\lambda_i + \lambda_{SO}) - 0.3(0.9)\lambda_T \lambda_{SO} \lambda_{Th}}$$

$$+ \frac{U_T^i (70000 - M_{Th})}{70000} . \quad (2.71)$$

where M_{Th} is the mass of thyroid.

3.0 DOCUMENTATION FOR "DOSE FORTRAN"

3.1 OBJECTIVE

"DOSE FORTRAN" is a software package, written in FORTRAN-77, that implements the methods expounded in Part 1 of Publication 30 of the International Commission on Radiological Protection (ICRP) to calculate committed dose equivalents from an internal radionuclide to organs and tissues of an adult "reference man" (as described in ICRP Publication 23). The program can consider any of the three major modes of intake of a radionuclide, namely, ingestion, inhalation, or submersion in a cloud of inert radioactive gas or elemental tritium. Specifically, the program can calculate the following:

- i) Specific committed dose equivalent HFIFTY (Sv/Bq) in 19 target organs,
- ii) Weighted committed dose equivalent WDOSE (Sv/Bq) in selected target organs,
- iii) Annual limit of intake ALI (Bq) of the radionuclide,
- iv) Derived air concentration DAC (Bq/m³) of the radionuclide in case of inhalation or submersion, and
- v) Specific effective energies in a table for 17 source and 19 target organs.

A list of the source and target organs along with the masses in reference man is given in Table 3.1.

TABLE 3.1 Masses of organs and tissues of "Reference Man" in this program.

No.	Source Organs	Mass (g)	No.	Target Organs	Mass (g)
1.	Bladder content	200.00	1.	Lungs	999.00
2.	Stomach content	250.00	2.	Thyroid	19.60
3.	SI content	400.00	3.	Testes	37.10
4.	ULI content	220.00	4.	Ovaries	8.27
5.	LLI content	135.00	5.	Red marrow	1500.00
6.	Kidneys	310.00	6.	Stomach wall	150.00
7.	Liver	1800.00	7.	SI + contents	1040.00
8.	Lungs	1000.00	8.	ULI wall	209.00
9.	Muscle	48200.00	9.	LLI wall	160.00
10.	Ovaries	11.00	10.	Liver	1810.00
11.	Pancreas	100.00	11.	Kidneys	284.00
12.	Trabecular Bone	1000.00	12.	Bladder wall	45.10
13.	Skin	2600.00	13.	Muscle	48200.00
14.	Spleen	180.00	14.	Bone surface cells	10500.00
15.	Testes	35.00	15.	Skin	2830.00
16.	Thyroid	20.00	16.	Spleen	174.00
17.	Total body	70000.00	17.	Uterus	65.40
18.	Cortical Bone	4000.00	18.	Pancreas	60.30
			19.	Total body	69900.00

3.2 REQUIREMENTS FOR EXECUTION

Some text files of auxiliary subroutines need to be linked to the central program "DOSE FORTRAN" for its execution. The subroutines (explained later) are as follows:

- i) DECAY1 FORTRAN,
- ii) ATOMNO FORTRAN,
- iii) ICLASS FORTRAN in case of inhalation,
- iv) F1VALU FORTRAN,
- v) FACTOR FORTRAN,

- vi) ICRP FORTRAN,
- vii) INGEST, INHALE, OR SUBMER FORTRAN,
- viii) PCLASS FORTRAN in case of inhalation,
- ix) RESPIR FORTRAN in case of inhalation,
- x) DECAY FORTRAN,
- xi) FRAC FORTRAN,
- xii) THALF FORTRAN,
- xiii) REFMAN FORTRAN,
- xiv) TFRAC FORTRAN,
- xv) TRNSFM FORTRAN,
- xvi) SPEFF FORTRAN,
- xvii) YERROR FORTRAN,
- xviii) INTRPT FORTRAN,
- xix) ENERGY FORTRAN,
- xx) I1 FORTRAN,
- xxi) SOURCE FORTRAN,
- xxii) UXP FORTRAN,
- xxiii) RESULT FORTRAN, and
- xxiv) FLOW FORTRAN

The directly accessed data files for decay schemes of radionuclides, number of transformations for radioactive alkaline earths, dose equivalent rates in body tissues from submersion, absorbed fractions of photon energies in organs, retention fractions of nuclides

in source organs, and the fractional transfer of the nuclides to the body fluid compartment are:

- i) ISOTIPS FILE (sequentially accessed),
- ii) ISOTOPE FILE,
- iii) ALPHA FILE,
- iv) BETA FILE,
- v) ELECTRN FILE,
- vi) DAUTER FILE,
- vii) POSITRN FILE,
- viii) ABSFRAC FILE,
- ix) RETENT FILE,
- x) BFFRAC FILE,
- xi) INDEXI FILE,
- xii) INDEXO FILE,
- xiii) EXCEPT FILE,
- xiv) LIST FILE, and
- xv) NOBLE FILE

3.3 EXECUTION OF THE PROGRAM

This program was written on the IBM Mainframe computer at Kansas State University. Conversational Monitor System (CMS) is the operating system used there to run under global control program (CP) which handles the resources of the mainframe computer. A feature of the CMS is the EXEC processor. A CMS EXEC processor is a CMS file that

contains executable statements. Hence, in this case, the "DOSE EXEC" file contains statements to retrieve, expand, and load the compiled text files of all supporting subroutines and the main program, thus functioning as a catalogued procedure for execution of the program. With the file on "A" (temporary) disk, one can invoke its execution by entering the word "DOSE".

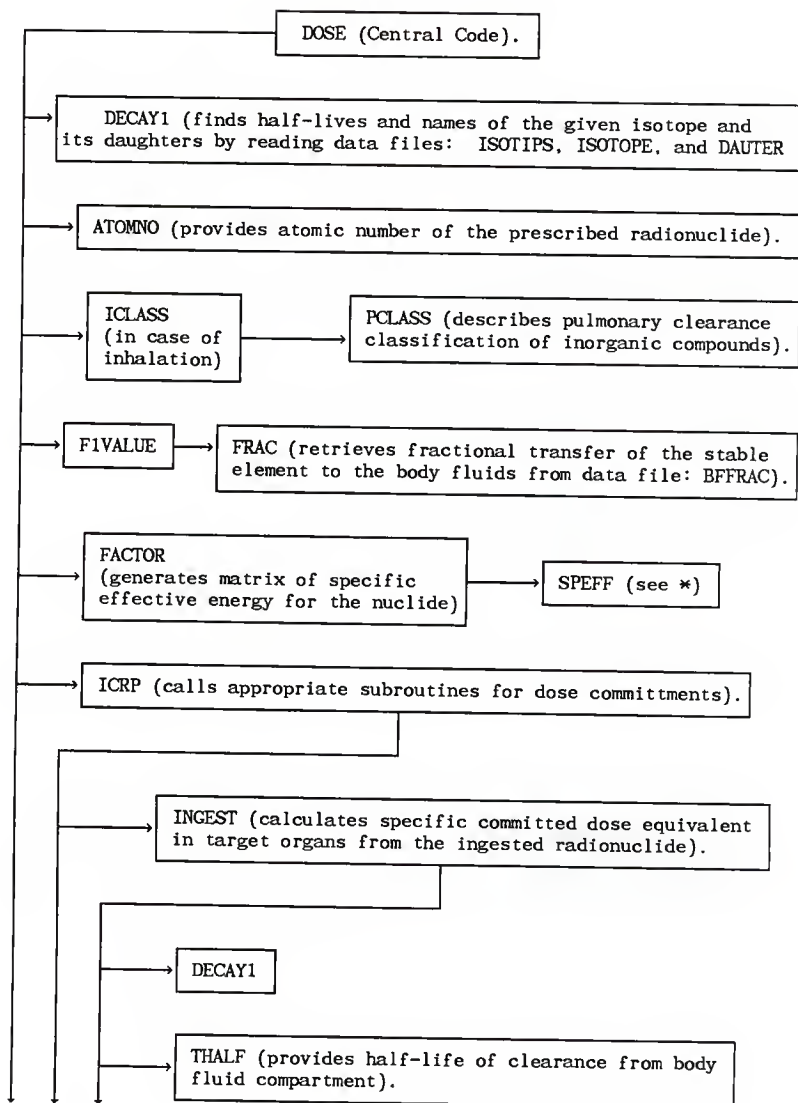
The main program in this package is "DOSE FORTRAN" which calls other subroutines to perform certain calculations. What follows are descriptions of the function and role of each subroutine as it is addressed by the calling program in execution. All three modes of exposure are considered. A summary of the program flow is illustrated in Fig. 3.1. We begin with the central code.

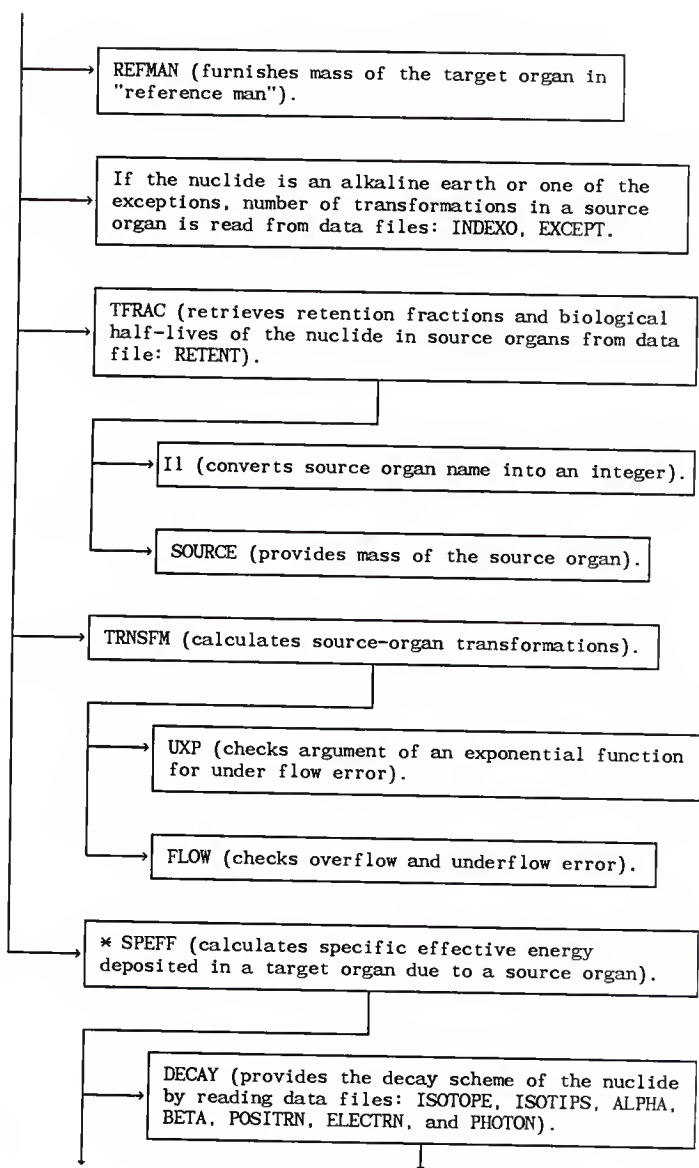
3.4 DOSE FORTRAN

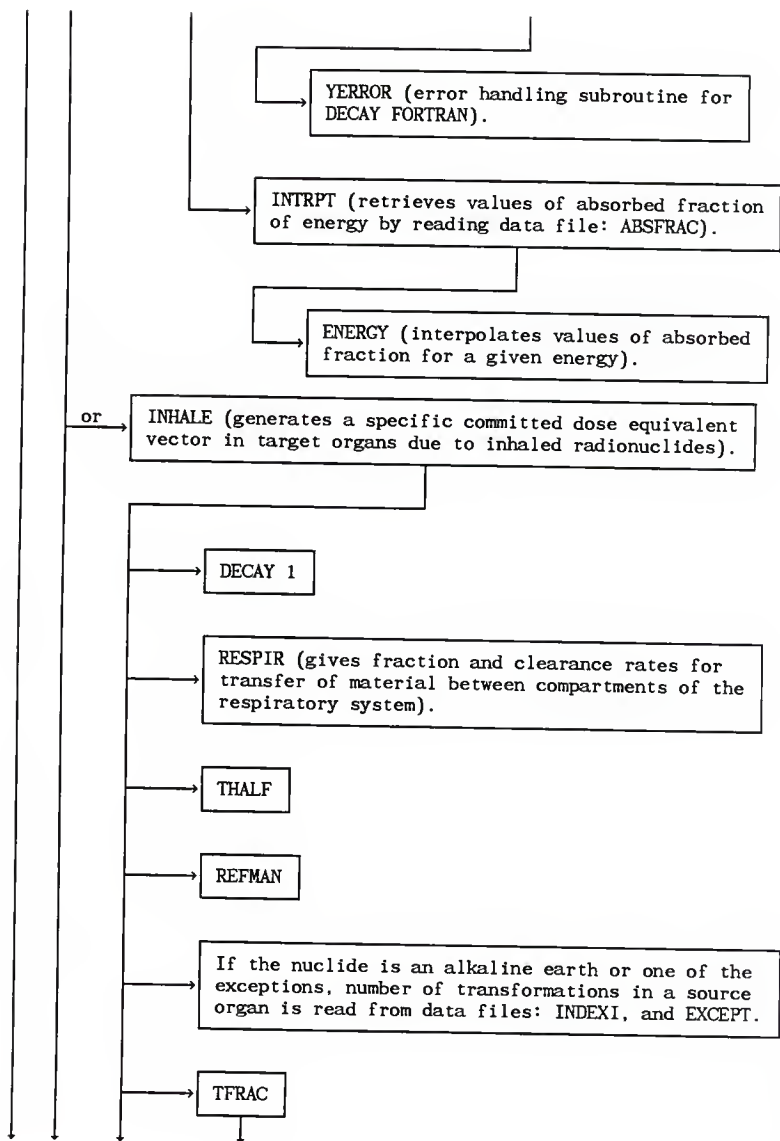
This is an interactive, user-friendly code which prompts the user to enter some basic information for its execution. To avoid confusion, an explanation of the screen-by-screen sequence in actual execution of the program is given below:

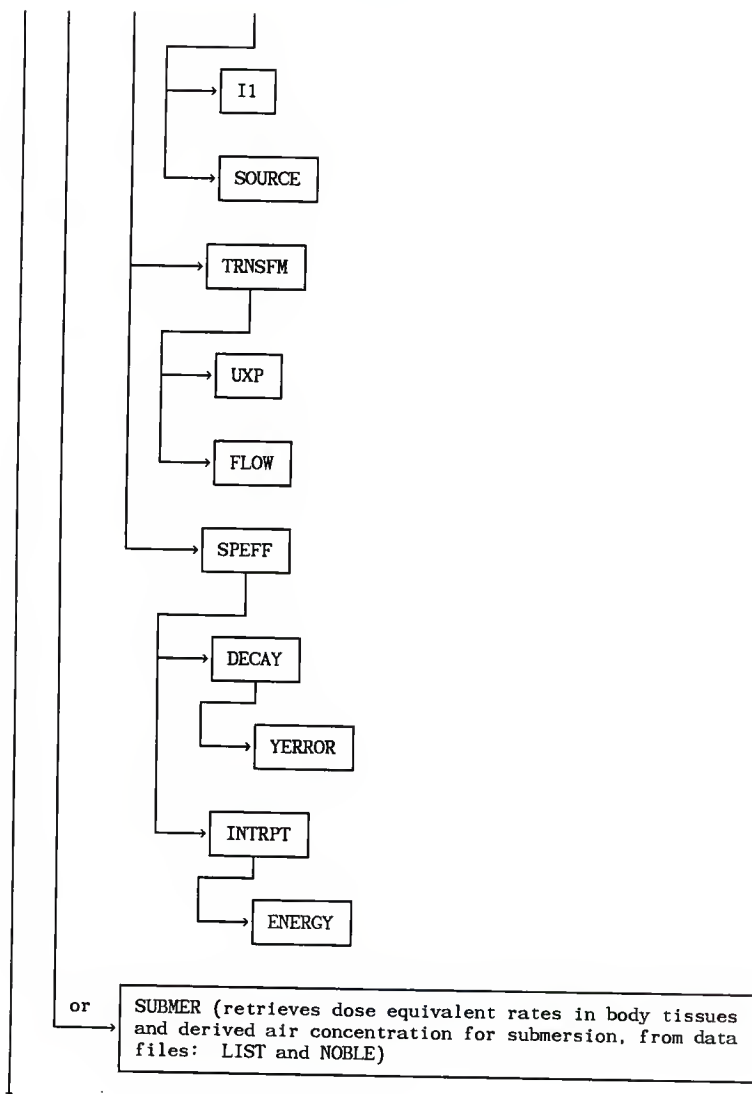
i) First and Second Screen: The first screen orients the user with the name of the program as well as the author, with the time and place of documentation of the code. The second screen explains the scope of the program.

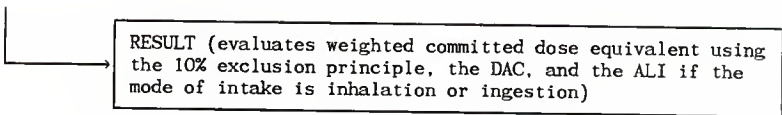
ii) Third Screen: There are three ways to provide the preliminary data required for a calculation. They are: 1) data input from the keyboard, 2) data input from a file, or 3) preparation of a











RESULT (evaluates weighted committed dose equivalent using the 10% exclusion principle, the DAC, and the ALI if the mode of intake is inhalation or ingestion)

FIG. 3.1. Points of call of subroutines in execution of the program.

data input file. The user is queried in this screen to opt for one of three choices. The choice is stored as an integer variable, NDATA which may have a value of 0, 1, or 2. Under choices 1 or 2, an error in program execution results in program termination.

If the user selects the option of data input from an already prepared file (NDATA = 1) residing on the temporary disk (A disk), the fourth screen inquires the name of the input file. The name is saved as a character variable FFILE(3). Since the CMS environment allows only 7 characters for the name of a file, a longer name is therefore truncated to 7 characters. The file is then opened and, according to the data read, the appropriate course of action is taken.

Also, if the user wishes to prepare a data input file, (NDATA=2) the fourth screen asks for a name that can be associated to such a file [FFILE(3)] and subsequent screens, then prompt the user for data from the keyboard which is then entered in the file. These screens for input data collection are also displayed in the event the user chooses the first option (NDATA=0) to enter data directly from the keyboard. These screens are explained below.

iii) Fourth screen: The user is prompted to enter a radionuclide identification according to displayed example of cesium-137 as CS-137. This is stored as the character variable WORD. If there is a mistake in the format of entry, the screen is displayed again until the entry is corrected. The screen also informs the user, that to begin termination of the program at this point, the return key may be pressed. When the return key is pressed, the user is questioned again

to confirm if termination is really what is desired. Of course, an assent results in the termination and a dissent shows the fourth screen again.

At this point, with the nuclide identified, a subroutine DECAY1 is called to find the daughters of the given radionuclide.

3.4.1 DECAY1 FORTRAN:

The objective of this subroutine is to access the half-lives and names of the daughters of the radionuclide under consideration. The data files used for this purpose are "ISOTIPS", "ISOTOPE", and DAUTER". The arguments of the subroutine are:

- WORD - Identification of the entered nuclide.
- RHALF - This is a vector of maximum length 50. It stores half lives of the given radioisotope and its daughters in sequential order of decay.
- ULIFE - This is a character vector of length 50. It saves units of half-lives of nuclides in RHALF as a character, e.g., S, M, H, D, or Y for seconds, minutes, hours, days, and years respectively.
- BRA - This is a vector of branching ratios of the given isotope (BRA(1)=1), and its daughters. Its maximum length is 50.
- RADIO - This is a vector of length 50. Its elements are characters, each of length 8. It stores identities (i.e., symbol-atomic weight, e.g., CS-137) of given nuclide and its daughter.

NO - This is an integer which stands for the number of daughters plus one (for the given isotope).

The given identification WORD of the nuclide is used to find a match in the sequentially accessed data file "ISOTIPS". Once a match occurs, a pointer or record number is read, which is then used to directly access the decay scheme from data file "ISOTOPE". The variables read are:

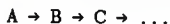
ERT - Name of the isotope.
JO - Atomic weight of the isotope.
J - Atomic number of the isotope.
B - Half life of the isotope.
U - Half life units (S, M, H, D, or Y).
K - Number of radioactive daughters.
L - Pointer or record no. for first daughter.
M - Number of alpha particles.
N - Pointer or record no. of first alpha particle.
I1 - Number of beta particles.
I2 - Pointer to first beta-particle record.
I3 - Number of positrons.
I4 - Pointer to first positron record.
I5 - Number of electrons.
I6 - Pointer to first electron record.
I7 - Number of photons.
I8 - Pointer to first photon record.

The pointer to the first daughter is used to directly access the branching ratio of the daughter from the data file "DAUTER". The variables read are:

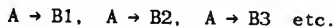
I9 - Pointer to daughter isotope. This record number could now be used to access the decay scheme of the daughter from "ISOTOPE", and so on.

YIELD - Branching ratio of the daughter.

The decay mode considered in this program is



Hence, if a nuclide decays into more than one daughter, that is, if



then the pointer and yield of only the daughter with the highest branching ratio is saved.

An inability to read any of the data files results in display of an appropriate error message and return to the fourth screen of the main program.

The half lives of the parent and the radioactive progeny are first converted into days if they are in any other units. If a daughter is a radioactive noble gas, then the daughter and its subsequent progeny are neglected and assumed to escape from the body. If the ancestral nuclide entered in the fourth screen is a noble gas, then an integer variable OPT is assigned a value of 3, and the following queries are skipped to the output screen, which asks for the output file name.

iv) Fifth Screen: The user is prompted to enter the sex of the exposed subject as either M or F. Character variable SEX stores the entry. An incorrect entry will again lead to same screen display until it is corrected.

v) Sixth Screen: The user is questioned whether or not, a full table of specific effective energies (S-matrix) of the nuclide for 17 source and 19 target organs is desired. To answer yes or no, the user is asked to enter the integer 1 or 2 respectively. Of course, an error is again handled with repeated displays of the same screen until proper entry is made.

If the user opts to see the table, the following screens are shown:

vi) Seventh Screen: It asks for the file name to which the matrix can be written. This name is stored as a character variable FFILE(1).

vii) Eight Screen: There are three units, i.e., MeV/g, rad/micro Ci.h, and mSv/GBq.h in which the S-matrix can be seen. The user is asked to choose one of them as an integer variable NU with a value of 1, 2, or 3.

If from the earlier call of subroutine "DECAY1" it is found that the radionuclide has daughters, then the ninth screen is as follows:

viii) Ninth Screen: The daughters of the nuclide are shown, and the user is prompted to enter integer 1 or 2 expressing yes or no to whether the S-tables of the daughters are desired. This expression is saved as an integer variable MON.

ix) Tenth Screen: At this point, the user is asked to choose one of the following options:

- 1) Continue data entry for calculation of dose commitments,
- 2) Conclude data entry and STOP, or

- 3) Proceed with calculations of S-matrix only.

Another choice is added if the user chooses the third option of preparation of a data input file in the third screen. It is:

- 4) Continue data entry for calculation of S-matrix only.

The choice is saved as an integer variable INRE with a value of 1, 2, 3, or 4 respectively. Of course, selection of option (1) here, would result in end of the program. If the user chooses to enter data from the keyboard in the third screen, and option number (3) here, then the query screens end, and the program proceeds with calculation of the S-matrix. However, if the choice was preparation of data input file in the third screen, then a selection of option (3) or (4) here, would first lead to writing of the data obtained from screen queries onto the file named by the user. After that, option (4) would display the fourth screen again, continuing the data entry process, while option (3) would read the written data and advance with calculation of the S-matrix, informing the user with a display of which calculation is being performed.

Now, regardless of the choice of either data entry from keyboard or preparation of an input file in the third screen, if option (1) is selected here, then the screen query continues as follows:

x) Eleventh Screen: The user is prompted to choose a mode of intake of the radionuclide by entering either 1 or 2 for ingestion or inhalation respectively. Integer variable OPT stores the selection.

xi) Twelfth Screen: The user is inquired if the transformations of the nuclide in source organs are fancied. The response can be

expressed as 1 or 2 for yes or no which is stored as integer variable MAIS. If the answer is yes, then the following two screens would ask for the name of a file, saved as character variable FFILE(4), to which the transformations can be written, and whether the transformations of the daughters (if any) are also desired. This response is stored as an integer variable ISAY which has a value of 1 or 2 for yes or no respectively.

3.4.2 ATOMNO FORTRAN:

At this point, this subroutine is called to assign an atomic number to the given radionuclide by comparing its symbol to the 103 in the program. A match results in an assignment of atomic number, or else, the user is asked through an error message to check the symbol and try again. The arguments of this subroutine are:

- SYM - First two characters of the variable WORD entered by the user which describes the symbol of the radionuclide. If, for instance in the case of phosphorus, the symbol is represented by only one character "P", then the second character is "-".
- KZ - This is the assigned atomic number by the subroutine and is naturally, an integer variable.

If the mode of intake was chosen to be inhalation in the eleventh screen, then the following subroutine is called.

3.4.3 ICLASS FORTRAN:

This subroutine provides the inhalation class of the given radionuclide. It first displays a laconic definition of the three classes, D, W, and Y, and then, questions the user if a detailed explanation aiding in selection of the pulmonary clearance classification is wished. If the user acquiesces, then another subroutine "PCLASS FORTRAN" is called, and after that the user is prompted to enter the inhalation class of the given radionuclide, which is the argument CLASS of the subroutine.

3.4.4 PCLASS FORTRAN:

This is an information file. It displays pulmonary clearance categories of different inorganic compounds which may aid the user in determining the inhalation class of the radionuclide. Its argument IWISH is merely the selection integer 1 indicating acceptance to view the file.

The next step is the determination of fractional transfer of the element from the GI system to the body fluids. For this purpose, subroutine FIVALU is called.

3.4.5 FIVALU FORTRAN:

An auxiliary function subprogram FRAC FORTRAN is required. The arguments of this subroutine are:

- KZ - Integer variable describing the atomic number of the given radionuclide, and
- F1 - The fractional transfer of the stable element from the GI system to the body fluids.

3.4.6. FRAC FORTRAN:

This function subprogram retrieves the fractional transfer F1 of the stable element to the body fluid compartment from data file "BFFRAC". Since the fraction may differ according to the inhalation class of the nuclide, the user is asked to enter the appropriate value from choices displayed on screen. Of course, a mistake in reading of data file is handled through an error message, suggested corrective action, and return to the fourth screen. The arguments of this subroutine are the atomic number of the nuclide KZ and the integer ITRACK, which is given a positive value in case of an error.

Again, if the mode of intake was chosen to be inhalation, the following screen is flashed.

xii) AMAD Screen: The user is inquired if the aerodynamic diameter (AMAD) for inhalation, is 1 micrometer. An affirmation or negation assigns a value of 1 or 2 respectively to integer variable LOT. In the event of a negative response, the user is prompted to enter the value of AMAD in micrometers between 0.1 and 20, which is stored as variable AMAD.

With these basic input values, the execution of "DOSE FORTRAN" can be initiated with call to appropriate subroutines. But before that, the name for output table for dose commitment results is needed.

xiii) Output Screen: A filename for output table is requested and a character variable FFILE(2) is assigned the name.

If the parent nuclide is a radioactive noble gas then at this point, the subroutine "ICRP" is called, and the following steps are ignored.

If the user is preparing a data input file, then at this stage a selection must be made from the following three choices:

- 1) Conclude data entry and STOP,
- 2) Continue data entry, or
- 3) Proceed with calculations.

Integer variable MORE is assigned a value of 0, 1, or 2 respectively according to choices described above. Except for option (1), choice of any other option would lead to writing of the data onto the named file. After that, option (2) would return to the fourth screen while option (3) would read the written data and progress with execution of "DOSE FORTRAN". To facilitate comprehension of the central code's working, most of the subroutines are explained in sequence of calling.

The step première in execution is S-matrix generation and printing of its results if, a positive response was given in sixth screen to view the S-table. A loop is initiated to do the same for the daughters if there are any, and if the user wishes to see their S-tables too.

3.4.7. FACTOR FORTRAN:

As mentioned earlier, this subroutine is called if a full specific effective energy (SEE) table (S-matrix) is requested in the sixth screen prompt of the main program. While this subroutine is generating the matrix, the main program flashes the message that it is calculating the S-matrix of the radionuclide transferred as an argument, and that the results are stored in the file named by the user. The arguments of this subroutine are:

- WORD - This is a character variable of length 8 which describes the parent radionuclide or its daughters found by the call of DECAY1 in the main program.
- NU - This is also a character variable. It has a length of 16 bytes, and it particularizes the choice of the user to see the SEE table in any of three sets of units, namely MeV/g, rad/micro Ci.h, or mSv/GBq.h.
- SFACT - This is the matrix of SEE values. Its size is 19 target organs in one dimension and 18 source organs in the other.
- NDATA - This is an integer variable which represents the option chosen in the main program of entering the data directly from keyboard or preparing a data input file.
- NUCLID - This character variable of length 8 identifies the parent radionuclide as entered by the user in the main program.
- PLIFE - This variable represents the half-life of the parent radionuclide converted in units of days.

The SFACT matrix is generated with uses of subroutines such as "DECAY FORTRAN" for the decay scheme of the radionuclide, and the subroutine "SPEFF FORTRAN", which in turn, calls other subroutines for effective energy absorbed in a tissue or organ. A detailed account of these subroutines follows later. The variable NDATA aids in properly redirecting the subroutine to the fourth screen in the main program if it has a value equal to zero or stopping the program if it has a value equal to one, in case of an error.

After generation of SEE values, stored in matrix SFACT, the main program writes these values as a table in the file FFILE(1) named by the user. If the mode of entry of data was keyboard, then these results are also written on screen for user's convenience.

At this juncture, if the user has prepared a data input file or had one on the A disk before the commencement of the program, i.e., if NDATA has a value of either 1 or 2 and, if a selection was made to proceed with calculations (only the S-matrix, INRE=3) in the screen prompt number 10, then the program will end here. However, if variable INRE has a value of 4 representing the wish to generate S-matrices of other nuclides as well, then the program would return to reading the data in the input file and will proceed according to the input. On the other hand, if the user were using the keyboard as mode of data entry, i.e., NDATA = 0, and if INRE = 3, the program would not end but could revert to the fourth screen allowing the user to end it personally or continue for any other nuclide.

Excluding the redirections in the above cases, the program will advance to calculate the dose commitments by calling the subroutine "ICRP FORTRAN".

3.4.8 ICRP FORTRAN:

This subroutine calls other appropriate subroutines for dose commitments according to the mode of intake. The arguments of this subroutine are:

- INTAKE - Integer variable with a value of either 1, 2, or 3 representing the mode of intake as ingestion, inhalation, or submersion respectively.
- WORD - Name of the given isotope, e.g., IN-113M a character variable.
- SEX - Sex of the subject as either M or F, a character variable.
- F1 - Fractional transfer of stable element from the GI system to the body fluids, a real variable.
- CLASS - In case of inhalation, pulmonary uptake classification of the nuclide, a character variable.
- AMAD - This is a real variable describing the activity median aerodynamic diameter of the nuclide in case of inhalation.
- ROB - Mass of whole body (70000 g) minus the masses of those organs and tissues mentioned in the metabolic model of ICRP-30 for a particular radionuclide.
- KZ - Atomic number of the nuclide.

- HFIFTY - Specific committed dose equivalent to target organ or tissue. This is a vector of length 24.
- US(I,J) - Matrix of transformations of nuclide I in source organ J.
- NDATA - An integer variable describing the options chosen in the main program of mode of data entry.
- DER - In case of submersion, this real variable represents derived air concentration for an inert radioactive gas.
- RISK - This real variable describes derived air concentration for an inert radioactive gas if it is determined by consideration of non-stochastic effects in case of submersion.
- ORGAN - This integer variable gives the organ or tissue number when the derived air concentration is determined by non-stochastic effects in case of submersion.

This subroutine merely calls subroutines "INGEST", "INHALE", or "SUBMER" according to the mode of intake transferred in the argument INTAKE as 1, 2, or 3 respectively. However, in case of inhalation, if the AMAD is not equal to 1 micrometer, then it also calculates the specific committed dose corresponding to the given AMAD from the 1 micrometer specific committed dose by law of proportions. The variable NDATA helps in properly re-routing the subroutine in case of an error. The three important subroutines mentioned above are described below.

3.4.9 INHALE FORTRAN:

This subroutine is called if the user chooses inhalation as the mode of intake. Its purpose is to generate a vector of specific

committed dose equivalents (Sv/Bq) in target organs or tissues due to inhaled radionuclides. The arguments of this subroutine are:

- WORD - This is a character variable of length 8 which identifies the given radionuclide.
- KZ - The atomic number of the nuclide.
- SEX - This is a character variable of length 1, and is either M or F for male or female respectively.
- HFIFTY(24) - This is a vector of length 24. The 19 elements of this vector represent the specific committed dose equivalents in 19 target organs or tissues.
- FNP(20) - This is a vector of length 20. The 19 elements represent the fractions of the committed dose equivalent in the reference target tissue resulting from deposition in the naso-pharyngeal (N-P) compartment of the lung model.
- FTB(20) - The 19 elements of this 20 element vector represent the fractions of committed dose equivalent in the target tissue resulting from deposition in the tracheo-bronchial compartment (T-B) of the model.
- FP(20) - The fractions of committed dose equivalent in the target tissue resulting from deposition in the pulmonary (P) region of the lung model are the first 19 elements of this 20 element vector.
- ROB - This is a variable which represents the mass of whole body (70000 g) minus the masses of those organs and tissues mentioned in the metabolic model of ICRP-30 for a

particular radionuclide. These organs are named in the data file "RETENT".

CLASS - This is a character variable of length 1 representing the pulmonary uptake classification of the radionuclide.

F1 - Fractional transfer of the stable element from the GI system to the body fluids.

US(I,J) - Matrix of transformations of nuclide I in source organ J.

A flowchart of this subroutine is shown in Fig. 3.2. This subroutine is initiated by calling the subroutine "DECAY1" to find the daughters of the radionuclide. Details of this subroutine were given earlier. After accession of the decay scheme from the above subroutine, half lives of parent and daughters, if not in units of days, are converted into days and then radiological constants are calculated. If the daughter is a radioactive inert gas, then that nuclide and its subsequent progeny are neglected and are assumed to escape out of the body.

With the use of the fraction of stable element F1 the fraction of the radioactive parent and its daughters to the body fluid compartment via the GI tract is calculated. This is saved in a vector of maximum length 50, named FBF.

The lung model of ICRP-30, described in a previous chapter, considers fractions of inhaled material to be deposited in three respiratory regions, the naso-pharyngeal passage (N-P), the trachea and bronchial tree (T-B), and the pulmonary region (P), the balance being the fraction exhaled. Initially, it is assumed that the activity

SUBROUTINE INHALE (WORK, KZ, SEX, CLASS,
F1, HFIFTY, FNP, FTB, FP, ROB, US, *)

CALL DECAY1 (WORD, RHALF, ULIFE, BRA, RADIO, NO, *12)
Purpose: Finds half-lives and names of the given
isotope and its daughters.

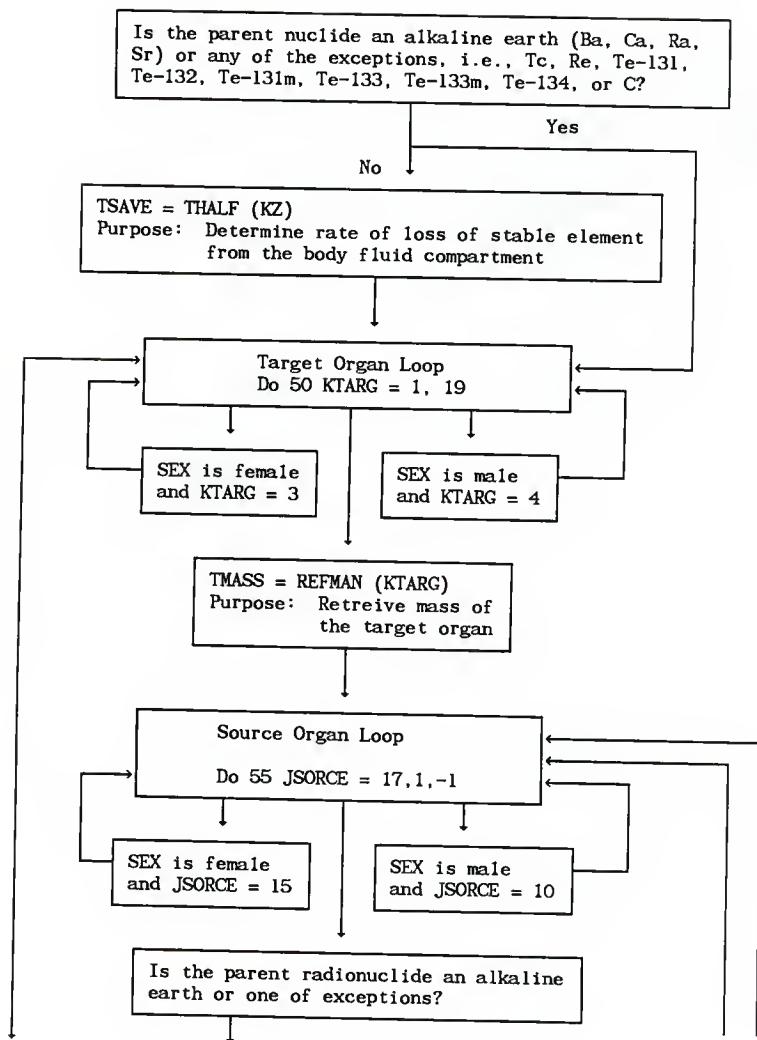
Convert half-lives into days and calculate radiological
constants of parent and its daughters.

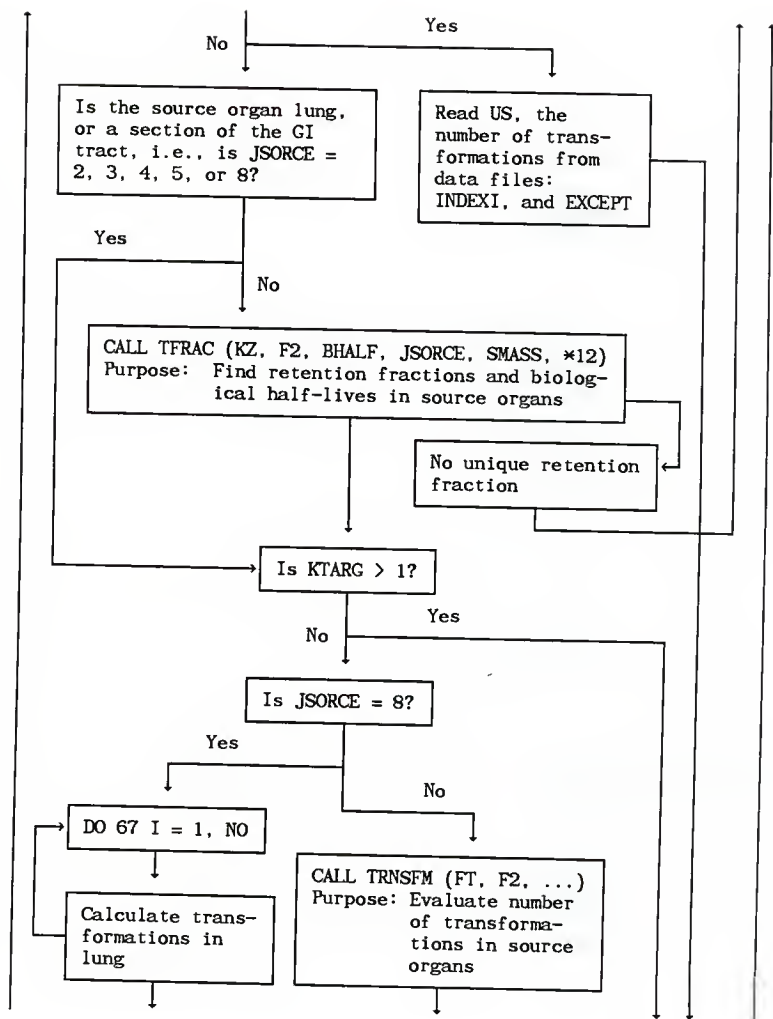
Evaluate fraction of inhaled radionuclide transferred
to the body fluid compartment via the GI tract, FBF.

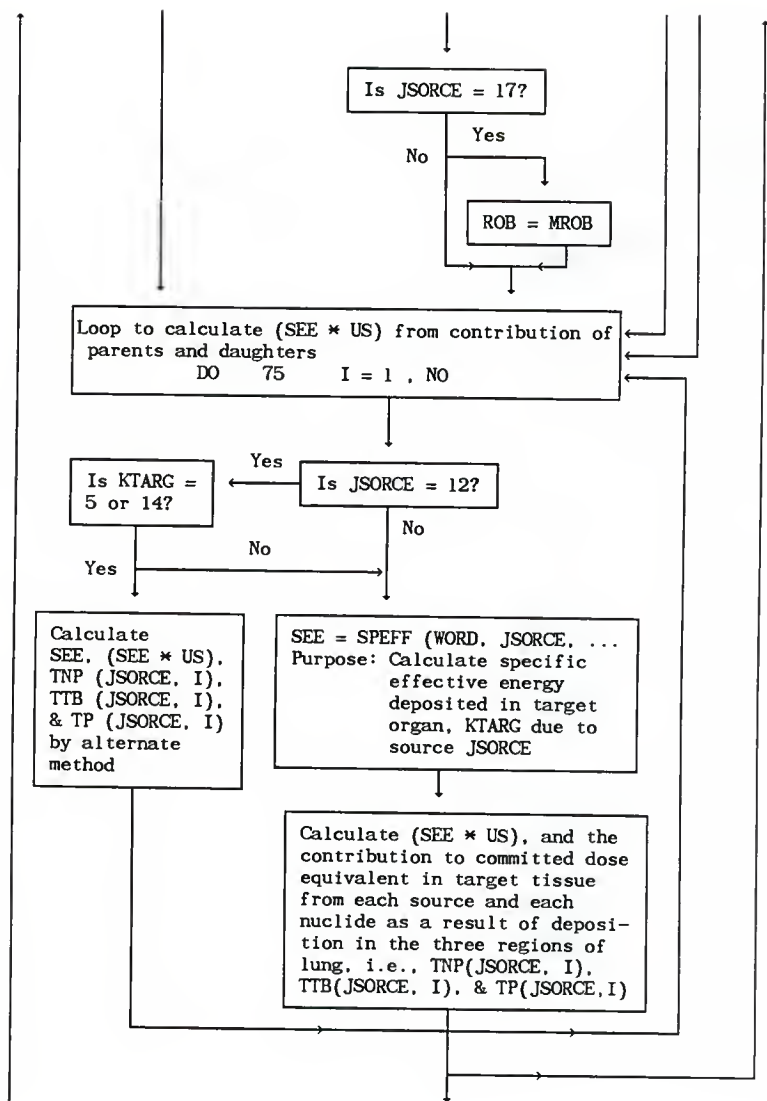
Assuming the AMAD = $1\mu\text{m}$, assign values to fractions
of inhaled material deposited in three respiratory
regions, the balance being the fraction exhaled.

CALL RESPIR (CLASS, FA, FB, FC, FD, ...)
Purpose: Retrieve fraction and clearance rates for
transfer between compartments of the lung.

Calculate transformations in each sub-compartment of
the lung. Use these results to find fraction of
parent and its daughters going directly to body fluids
FBFDIR, fraction going directly to GI tract FGI, and
total initial activity in the body fluid compartment
FT.







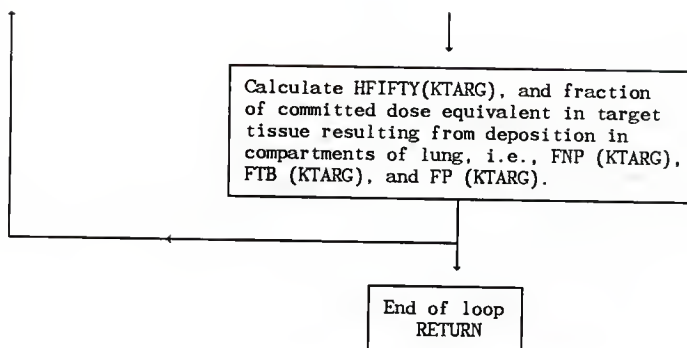


FIG. 3.2 Flowchart of "INHALE FORTRAN"

median aerodynamic diameter (AMAD) of the nuclide inhaled is 1 micrometer, and the corresponding deposition fractions in each compartment of the lung are utilized for calculation of specific committed dose equivalent. The reason for this assumption is that, even in alternate cases, when the AMAD is not 1 μm , values of HFIFTY for 1 μm AMAD are used to estimate HFIFTY in those cases, using the law of proportions.

The clearance rates and fractions in each sub-compartment of the three major divisions of the lung is first initialized as zero and then, a subroutine "RESPIR" is called for their values.

3.4.10 RESPIR FORTRAN:

This subroutine merely provides the fraction and clearance rates for transfer of nuclide between compartments of the lung according to the inhalation class of the nuclide. The arguments are:

- CLASS - Character variable of length 1 describing inhalation class of the nuclide.
- FA - Fraction of material deposited in the body fluid compartment from the nasal passage region (N-P).
- FB - Fraction deposited in the GI tract from the nasal passage.
- FC - Fraction deposited in the body fluid compartment from the trachea and bronchial tree region (T-B).
- FD - Fraction deposited in the GI tract from the trachea and bronchial tree region (T-B).

- FE - Fraction deposited in the body fluids from the pulmonary region (P).
- FF - Fraction transferred to the T-B region from the P region with a half-life.
- FG - Fraction transferred to the T-B region from the P region with a different half life.
- FH - Fraction transferred to the lymphatic system (L) from the P region.
- FI - Fraction deposited in the body fluids from the L region with a half life.
- CLA - Clearance rate of material from the N-P to the body fluids which is found from dividing $\ln 2$ by the removal half time (in days) in each compartment.
- CLB - Clearance rate of material from the N-P to the GI tract.
- CLC - Clearance rate of material from the T-B to the body fluids.
- CLD - Clearance rate of material from the T-B to the GI tract.
- CLE - Clearance rate of material from the P to the body fluids.
- CLF - Clearance rate of material from the P to the T-B region.
- CLG - Clearance rate of material from the P to the T-B region.
- CLH - Clearance rate of material from the P to the L region.
- CLI - Clearance rate of material from the L to the body fluids.
- CLJ - Clearance rate of material from the L to the body fluids.

With the use of these fractions and clearance rates, transformations in various sub-compartments of the lung for the parent and its daughters are assessed, and these transformations allow calculation of the fraction of the inhaled radionuclide and its daughters transferred directly to the body fluid compartment, and the fraction transferred to the GI tract. The former is saved as a vector FBFDIR, and the latter as a vector FGI. Both vectors are of maximum length 50. These results aid in determining the total initial activities of the given radionuclide and its daughters in the body fluid or transfer compartment. The activities are saved as a vector FT (length = 50).

Since different elements have different clearance times from the body fluid compartment, function subprogram "THALF" is called to determine the suitable half-life of clearance. However, if the parent nuclide is an alkaline earth (Ba, Ca, Ra, Sr) or one of the exception nuclides, i.e., Tc, Re, Te-131, Te-132, Te-131m, Te-133, Te-133m, Te-134, or C, this step is skipped to the target organ loop.

3.4.11 THALF FORTRAN:

This function subprogram uses the argument KZ, the atomic number of the nuclide, to provide the half-life of clearance from the transfer compartment. The daughters are assumed to have the same half-life of clearance as the parent.

With this half-life of clearance, the rate of loss of the stable element from the transfer compartment TOONST, is calculated. After that, the source-organ transformations in each source organ for the parent and its daughters are initialized to zero, and then, an outer loop to calculate HFIFTY in each target organ, is commenced. The target organs ovaries and testes are skipped if the sex of the subject is male or female respectively. To determine the mass of each target organ, a function subprogram named "REFMAN" is called.

3.4.12 REFMAN FORTRAN:

This function subprogram uses the target number as the argument and returns the mass as shown in Table 3.1.

Some variables which will be described later as they are used, are initialized to zero at this point. For each target organ in the outer loop, a nested loop to add contribution from all source organs is initiated. As in the target organ loop, if the subject is male, ovaries are omitted as source organ and similarly testes are ignored if the subject is female.

At this juncture, if the ancestral nuclide inhaled is an alkaline earth or one of the exceptions described above, the number of transformations in each source organ are not calculated but retrieved from data files. The first letter of the nuclide symbol is used to search for a match in file "INDEXI". Once a match occurs, the values of F1 (described above) and the class of the nuclide are compared.

Beside each value of F1 and class, is a record number, which is read on proper match. This record number is a pointer for file "EXCEPT". Using this record number, the numbers of transformations for different source organs are read from this file.

In the case of nuclides technetium (Tc) and rhenium (Re), an anomaly exists. The ICRP 30 describes transformations in stomach wall as well as stomach content as source organs. However, in this report both are summed and treated as only one source organ, the stomach content. Similarly, the ICRP-30 also gives specific effective energy values (described later) for both source organs. Since this program cannot generate specific affective energy values for stomach wall as source organ, the summed value of transformations is multiplied by the SEE values for stomach content as source organ. Although this treatment does not affect the results of dose committment in other target organs appreciably, it does lead to underestimation as compared to ICRP-30 in the case of target organ stomach wall and hence, must be used with caution.

For all other nuclides, to find the fractions of nuclide retained in source organs as a result of transfer from the body fluid compartment, a subroutine "TFRAC" is called. The contribution to committed dose in organs of the GI tract and lung through this route is minimal. Since the nuclide enters the body-fluid compartment after passage through these organs, the major contribution is direct. Moreover, this minimal part is taken care of in the source organ total body.

3.4.13 TFRAC FORTRAN:

The purpose of this subroutine is to retrieve the retention fractions F2 and the biological half-lives BHALF of the nuclide in the source organs from data file "RETENT". The auxiliary subroutines needed are "I1" and "SOURCE". The arguments used are:

- KZ - The atomic number of the nuclide.
- F2 - This describes the fraction retained in source organ except organs of the GI tract and lung. It is a vector of length 3 since source organs are modelled to have 3 compartments.
- BHALF - This vector of length 3 gives the biological half-lives of the fractions in the source organs.
- JSORCE - This is the source organ number as shown in Table 3.1.
- SMASS - This variable invokes the function subprogram "SOURCE" to obtain the mass of the source organ, except when the source organ is total body. In that case, if other organs are linked with retention fractions, then this variable is assigned a value of $70000 - (\sum_i M_i)$ where M_i are the masses of organs associated with different retention fractions.

A flow diagram of this subroutine is illustrated in Fig. 3.3. A loop is started to access retention fraction and biological half-lives associated with source organs for a given nuclide. Variable C2 describes the source organ name. Other variables D, E, and F describe the retention fractions in different compartments of source organ, and G, H, B read the corresponding biological half lives. After retrieving

the source organ name associated with retention fractions, and biological half-lives from the data file, function subprogram "I1" is called with the source organ name C2 in alpha numeric characters as argument. "I1" converts C2 to an integer from the source list presented in Table 3.1.

The source integer is now compared with JSORCE, the argument. If they are equal, then the corresponding retention fractions and biological half-lives associated with C2 are equated with F2 and BHALF respectively. The function subprogram SOURCE is called with source organ integer as argument. This function subprogram provides the mass of the source organ which is equated with variable SMASS. With these values, the subroutine "TFRAC" is returned.

If, however, the source integer linked with C2 does not match the JSORCE, then there are two possibilities. The obvious one is that the loop is continued until a match is found, and then the values returned, or no match found, and values = 0 are returned. The other route is that if JSORCE is 17, i.e., the source organ is total body and if the source integer corresponding to C2 is not 18, linked with "all other" then the masses of source organs are summed, and the loop continued, until the integer is 18, and at that time, retention fractions and biological half-lives related to it are taken to be the ones sought. The SMASS variable is, however, assigned a value of 70000 minus the previous sum of source organs, rather than 70000, the mass of total body. This mode of accession works because the records in the data

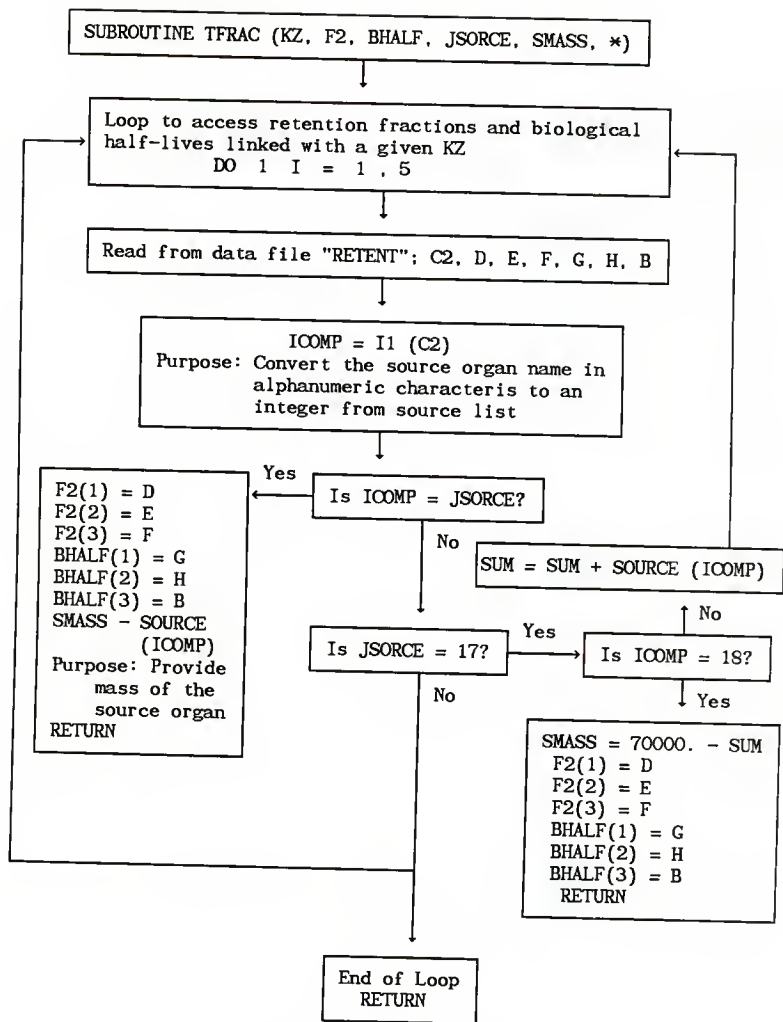


FIG. 3.3. Flowchart of Subroutine "TFRAC FORTRAN".

file are entered in such a way that for a given KZ, entry of "all other" is always at the end, after specific organs.

Of course, an error in reading or accessing a data file is disclosed by a display of appropriate message and suggested corrective action.

If a source organ does not have a unique retention fraction, that is, after the subroutine call, the retention fractions and biological half-lives for that organ are zero, then the source organ is skipped because often it will be included in the source organ total body. Since the transformations are evaluated in the source organ, their values remain the same for different target organs. Thus, if the target organ number is greater than 1 in the outer loop, repeated evaluation of source organ transformations is bypassed. Also if the source organ is lung, the subroutine for transformations is not called. Instead, the transformations calculated in each subcompartment earlier are summed, except for the N-P region. This treatment, as explained in the previous chapter, is based on the assumption that the dose received by the nasopharyngeal region for most particles sizes, is small in comparison with the doses received by other regions and hence, can be neglected. For all other sources, though, the subroutine "TRNSFM" is called.

3.4.14 TRNSFM FORTRAN:

This subroutine provides the source-organ transformations of the parent and its daughters in each source organ. The arguments are:

- FT - A vector of length NO (variable dimensioning) which describes the total initial activity of the given radionuclide and its daughters in the transfer compartment.
- F2 - A vector of length 3, which describes retention fractions in three compartments of the source organ.
- BHALF - Also a vector of length 3 which relates to the biological half-lives of the fraction retained in source organs.
- RCONST - This is a vector of length NO which describes the radiological constants of the parent and the daughters.
- NO - Integer variable describing number of daughters plus one for the parent.
- BRA - Branching ratio of the parent [BRA(1)=1] and its daughter saved as a vector of length NO.
- US - This is a matrix of size 20 x NO. The first dimension refers to the source integer number, and the second to the nuclide of interest. This array describes the transformations in a particular source organ for a particular nuclide.
- TCNST - Rate of loss of stable element from the transfer compartment.
- JSORCE - Source integer number.
- F1 - Fractional transfer of the stable element to the body fluid compartment.

- IProg - Integer variable identifying the calling subroutine as "INHALE" if value = 1, and "INGEST" if value = 0.
- FGI - A vector of length NO, describing the fraction of the inhaled radionuclide and its daughter transferred to the GI tract.
- SMASS - Variable describing mass of the source organ except if the source organ is total body and other organs of mass M_i have unique retention fractions. In that event, the variable equals $70000 - \sum_i M_i$.
- UROB - A vector of length NO describing the total number of transformations of each nuclide in rest of the body of mass, $70000 - \sum_i M_i$ where M_i is the mass of organ i for each unique retention fraction.
- MROB - This is a real variable which is assigned the value equal to $70000 - \sum_i M_i$ when the source organ is total body.
- KZ - Atomic number of the given radionuclide.

This subroutine begins with the calculation of biological constants BCONST from the biological half-lives in different compartments of the source organ. Depending on the mode of intake, i.e., the value of IProg, the calculations for organs of the GI tract are directed to appropriate line numbers for the evaluation of source-organ transformations by alternate methods.

An outer loop is initiated for the calculation of US for each nuclide. If TCONST is equal to zero, that is, there is instantaneous

transfer to the tissue compartment, then calculation of UTJ, the vector of length NO describing transformations of the nuclide under consideration in the transfer compartment, is omitted. Otherwise, a nested loop calculates UTJ for each nuclide. Another nested loop evaluates the transformations in the three compartments of the source organ, and adds them. Of course, the initial activity of the nuclide in the source organ after transfer from the body fluid compartment is first calculated within the loop. Transformation in each compartment of the source organ is stored as a variable UJ. Use of UTJ and UJ allows calculation of US(JSORCE, J) where J is the nuclide under consideration.

If the source organ is total body, then the source-organ transformations calculated are equated with UROB(J). This quantity is associated with transformation in "rest of the body" and the source-organ transformations used in conjunction with total body as the source organ is over estimated by multiplying by 70000/SMASS, where SMASS as explained previously is $70000 - \sum_i M_i$, M_i being the mass of the organ with unique retention fraction. This overestimation in case of total body is compensated for each organ with unique retention fraction by subtracting the quantity, $M_i \times \text{UROB}(J) / (70000 - \sum_i M_i)$ for organ i. The outer loop for each nuclide is closed at this point and the value of US(JSORCE,J) is returned.

As remarked earlier, for organs of the GI tract, depending on mode of intake, there are separate blocks for calculation of US(JSORCE,J), which follows the loop described above. In the case of iodine, a

separate three compartment model is described in the previous chapter. Transformations in the source organs are calculated using this model. A flow chart can be seen in Fig. 3.4.

After the determination of US, another nested loop is initiated to calculate the product of US and specific effective energy absorbed in the tissue for each nuclide. For this purpose, use of another function subprogram called "SPEFF" is required.

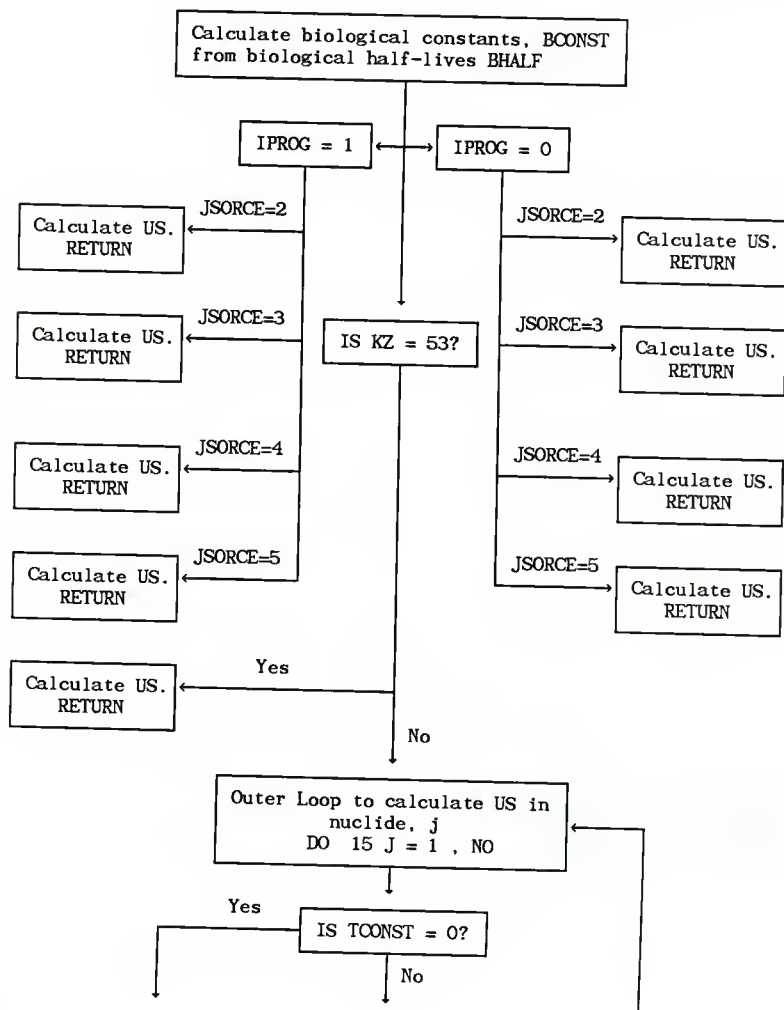
3.4.15 SPEFF FORTRAN:

This function subprogram calculates the specific effective energy deposited in each target organ due to each source organ. The arguments transferred are:

- WORD - Identification of the nuclide.
- JSORCE - Source integer number.
- KTARG - Target mass in grams.
- LOOP - Integer variable which has a value of either 0, 1, or 2.
- MOTS - Identification of the parent nuclide.
- PLIFE - Half life of the parent in days.

At the outset, the source number is compared with the target number according to Table 3.1. If the organs are the same, the integer variable ICOM is assigned a value of 0, otherwise a value of 1. For the decay scheme, a subroutine "DECAY" is called which will be explained later. The character variable SAVE stores the identification of the radionuclide when this function subprogram is first called, and

SUBROUTINE TRANSFM (FT, F2, BHALF, RCONST, NO, BRA, US, TCONST
JSORCE, F1, IPROG, FGI, SMAS, UROB, MROB, KZ)



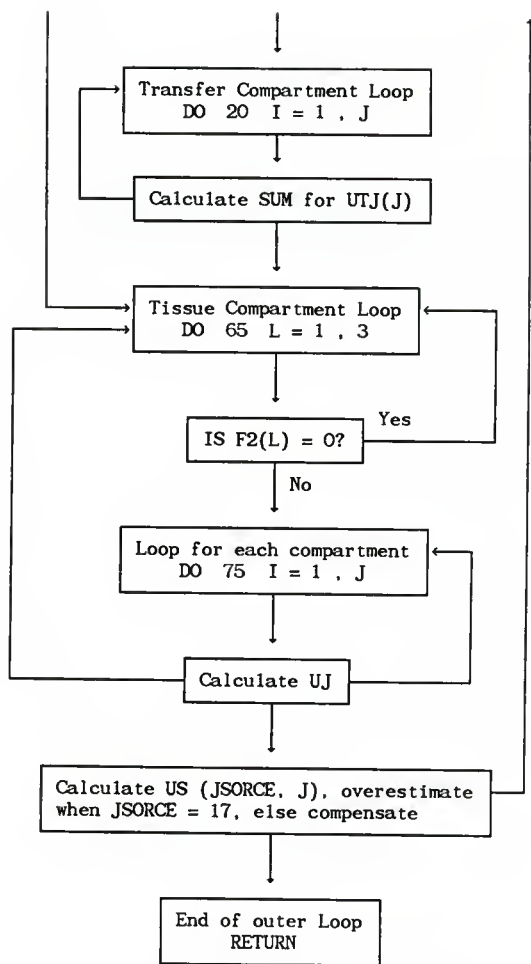


FIG. 3.4. Flowchart of "TRNSFM FORTRAN"

hence when it is equal to WORD, the function does not have to call "DECAY" again. At this point, if positrons are found in the decay scheme, corresponding annihilation photons are added to the list of decaying photons.

There are alternate methods for calculation of SPEFF for target organs of the GI tract, bladder, and bone. For all organs, there are five blocks for treating particles alpha, beta, positron, electron and photon. For target organs other than the GI tract, bladder, and bone, in the case of charged particles, the specific absorbed fraction in the target organ $\hat{A}F$ is zero if the source and target organs are not equal, except in the following cases:

- i) When the source organ is total body, the specific absorbed fraction equals $1/69900$.
- ii) When the target organ is total body and the source organ is bladder content, the specific absorbed fraction is equal to $45.1/(2 \times 200 \times 69900)$.
- iii) When the target organ is total body and the source organ is stomach content, the specific absorbed fraction equals $150/(2 \times 250 \times 69900)$.
- iv) When the target organ is total body and the source is SI content, $\hat{A}F = 640/(2 \times 400 \times 69900)$.
- v) When the target organ is total body, and the source organ is ULI content, $\hat{A}F = 210/(2 \times 220 \times 69900)$.
- vi) When the target organ is total body, and the source organ is LLI content, $\hat{A}F = 160/(2 \times 135 \times 69900)$.

vii) When the target organ is total body, and the source organ is any organ except ones described above, $\hat{AF} = 1/69900$.

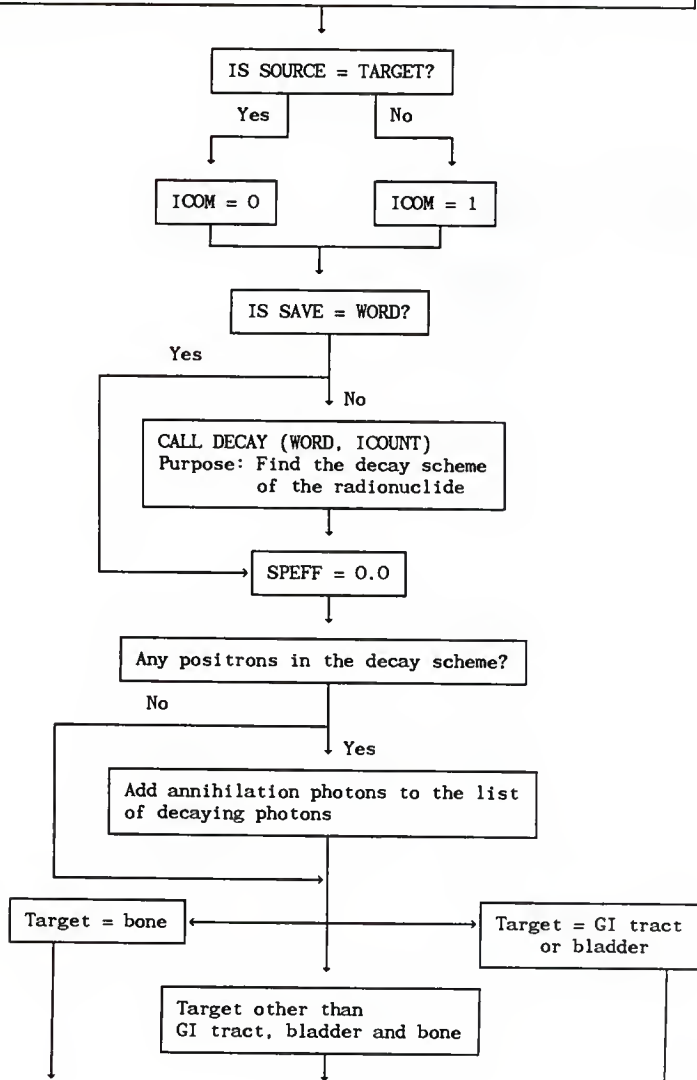
In the case of photons, the same cases hold true if the energy is less than 0.01 MeV. For higher energies a function subprogram "INTRPT" is called which will be explained later.

For target organs of the GI tract, bone and bladder, in the case of charged particles, the specific absorbed fraction is zero, if the sources and targets are not the same except when the source is total body. In that case, $\hat{AF} = 1/69900$. For photons, the same conditions hold true if the energy is less than 0.01 MeV, or else "INTRPT" is called. A flow diagram is sketched in Fig. 3.5.

3.4.16 DECAY FORTRAN:

This subroutine provides the decay scheme of the radionuclide. The data files are based on the radioactive decay data tables by D. C. Kocher, DOE/TIC-11026 (1981). It uses character variable WORD and integer variable ICOUNT as arguments. ICOUNT is normally 0, until an error occurs, in that case, its value is 1. Similar to "DECAY1", the symbol in WORD is used to find a match in the sequentially accessed data file "ISOTIPS". Once a match is found, a record number ITR is read which is then used to access directly the decay scheme from data file "ISOTOPE". Any error in this subroutine, is recorded as a particular element of an integer vector, for which subroutine "YERROR" is called, which will be explained later. Variables used to access "ISOTOPE" are:

SUBROUTINE SPEFF (WORD, JSORCE, KTARG, TMASS, LOOP, MOTS, PLIFE)



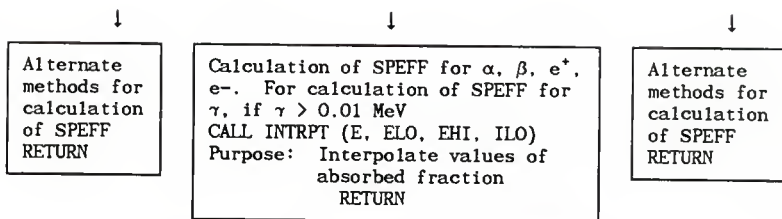


FIG. 3.5. Flowchart of Function Subprogram "SPEFF FORTRAN"

ERT - Name of the isotope.

JO - Atomic weight.

J - Atomic number.

B - Half-life.

C - Half-life units (S, M, H, D, Y).

K - Number of daughters.

L - Pointer to first daughter record.

M - Number of alpha particles.

N - Pointer to first alpha particle record.

I1 - Number of beta particles.

I2 - Pointer to first beta particle record.

I3 - Number of positrons.

I4 - Pointer to first positron record.

I5 - Number of electrons.

I6 - Pointer to first electron record.

I7 - Number of photons.

I8 - Pointer to first photon record.

The pointer to first alpha particle is used to access the data file "ALPHA". The variables read are:

AL1 - Energy in MeV.

AL2 - Intensity.

The pointer to beta is used to access the data file "BETA" with the variable being:

BE1 - Endpoint energy in MeV.

BE2 - Average energy in MeV.

BE3 - Intensity.

The pointer to positron is the key to access the data file "POSITRN" for reading the following variables:

POS1 - Endpoint energy in MeV.

POS2 - Average energy in MeV.

POS3 - Intensity.

The pointer to electron is the record number to access "ELECTRN".
Variables read are:

ELE1 - Energy in MeV.

ELE2 - Intensity.

Similarly, pointer to photon is used for file "PHOTON" with variables:

PHO1 - Energy in MeV.

PHO2 - Intensity.

Each of these variables starting from AL1 are stored as a vector for the decay of parent and daughters, and all of these vectors are common to function subprogram "SPEFF". Besides these vectors other variables that are common are M, I1, I3, I5, I7, HLIFE. HLIFE, a variable, describes the half lives in units of days.

3.4.17 YERROR FORTRAN:

This is the error handling subroutine for the subroutine "DECAY FORTRAN". It gives error messages when called from "DECAY" in case of an error in reading data files, discloses the source of error as "DECAY", and suggests appropriate action. The argument of this

subroutine is DECERR(7). This is an integer vector of length 7. Each kind of error relates to a particular element of this vector. The element is assigned an integer value greater than zero in case of an error.

3.4.18 INTRPT FORTRAN:

This function subprogram is called by "SPEFF" for calculation of SPEFF in the case of photon decay for energy greater than 0.01 MeV. The purpose of this subprogram is to interpolate the values of absorbed fraction. The data file required is "ABSFRAC" and the subroutine "ENERGY". The arguments of this subprogram are:

E - Energy of gamma.

JSORCE - Source integer.

KTARG - Target integer.

It utilizes subroutine "ENERGY" to obtain the upper and lower bounds of energy which aids in interpolation. The data file "ABSFRAC" is used to read absorbed fraction for the upper and lower bounds of energy.

After determining the specific effective energy SEE the product of SEE and US is required for calculation of HFIFTY from contributions of all radionuclides. If the source organ is mineral bone and the target is either red marrow or bone surface cells then the contribution from both trabecular bone and cortical bone as part of mineral bone is calculated by first calling "SPEFF" with a value of variable LOOP as 0

which implies photon decay from radionuclides, then with a value of LOOP = 1 which takes care of charged particle dose in trabecular bone, and finally with LOOP = 2 which accounts for charged particle dose in cortical bone.

For other source organs, value of LOOP = 0. To determine the contribution to committed dose equivalent in target tissue from each source and each nuclide as a result of deposition in the N-P, T-B and P regions of the lung model, three matrices named TNP, TTB, and TP are evaluated which have a maximum size of $20 \times NO$ where 20 represents each source integer, and NO, the number of nuclides.

The products (SEE \times US) are added from contribution of each source in variable GRNSUM. The source loop is now closed at this point. FNP, FTB, FP, and HFIFTY described earlier is calculated at this point for each target organ and the target loop is then closed. The calculated values are returned to the subroutine "ICRP".

If the value of AMAD is not equal to 1, the subroutine "ICRP" calculates the committed dose by law of proportions, and then returns the calculated value to the control code "DOSE".

As mentioned earlier, the subroutine "ICRP" will call the subroutine "INGEST" to calculate the specific committed dose equivalent in target organs from the ingested radionuclide, if the value of variable, INTAKE is 1. The mode of operation of this subroutine is very similar to "INHALE" and hence the subroutines called by "INGEST" which are common to "INHALE" are not described to avoid redundancy.

3.4.19 INGEST FORTRAN:

The arguments passed by the subroutine "ICRP" for calculation of the specific committed dose equivalent due to the ingested radionuclide are:

- WORD - The character variable of length 8 which identifies the radionuclide.
- KZ - The atomic number of the nuclide.
- SEX - Character variable of length 1 identifying the sex of the exposed subject.
- F1 - Fractional transfer of the stable element from the GI system to the body fluids.
- HFIFTY(24) - The first 19 elements of this vector represent the specific committed dose equivalents in 19 target organs or tissues.
- ROB - This variable stands for the mass of whole body minus the masses of those organs and tissues with unique retention fractions mentioned in the metabolic model.
- US(I,J) - This is a matrix of transformations of nuclide I in source J.

This subroutine like its counterpart "INHALE" is initiated by calling "DECAY1" for half-lives and names of the parent isotope and its daughters. The half-lives are converted into days and radiological constants calculated. Again, if the progeny is a radioactive inert gas, it is assumed to escape out of the body.

The initial activity FT of the parent and its daughters in the transfer compartment is evaluated using the fractional transfer of the stable element from the GI tract to the body fluids.

Except for the alkaline earths and the nuclides Tc, Re, Te-131, Te-131m, Te-132, Te-133, Te-133m, Te-134, and C, for which the number of transformations in source organs are retrieved from a data file "EXCEPT", the half life of clearance for other nuclides is obtained from function subprogram "THALF".

A target organ loop to calculate HFIFTY is started at this point. Mass of each target organ is given by function subprogram "REFMAN". For each target organ, contribution from all source organs is evaluated through a nested source loop.

To calculate the number of transformations in source organs, the retention fractions of the nuclides in these source organs must first be determined. Subroutine "TFRAC" retrieves these retention fractions and their biological half lives in the source organs. After this step, subroutine "TRNSFM" is called for the source organ transformations of each nuclide, and then subroutine "SPEFF" for specific effective energy in each target organ due to each source organ. The product of both the quantities is added from contribution of all sources in each target organ. The source loop is closed and then the specific committed dose equivalent in each target organ is calculated. After covering all 19 target organs, the loop is closed and the values returned to "ICRP" which in turn routes it to "DOSE". A flow diagram for "INGEST" is shown in Fig. 3.6.

SUBROUTINE INGEST (WORD, KZ, SEX, F1, HFIFTY, ROB, US, *)

CALL DECAY1 (WORD, RHALF, ULIFE, BRA, RADIO, NO, * 12)

Convert half-lives into days and calculate
radiological constants for parent and daughters

Is the parent nuclide an alkaline
earth or one of the exceptions?

Yes

No

Calculate total initial activity of each
nuclide in the transfer compartment, FT

TSAVE = THALF (KZ)

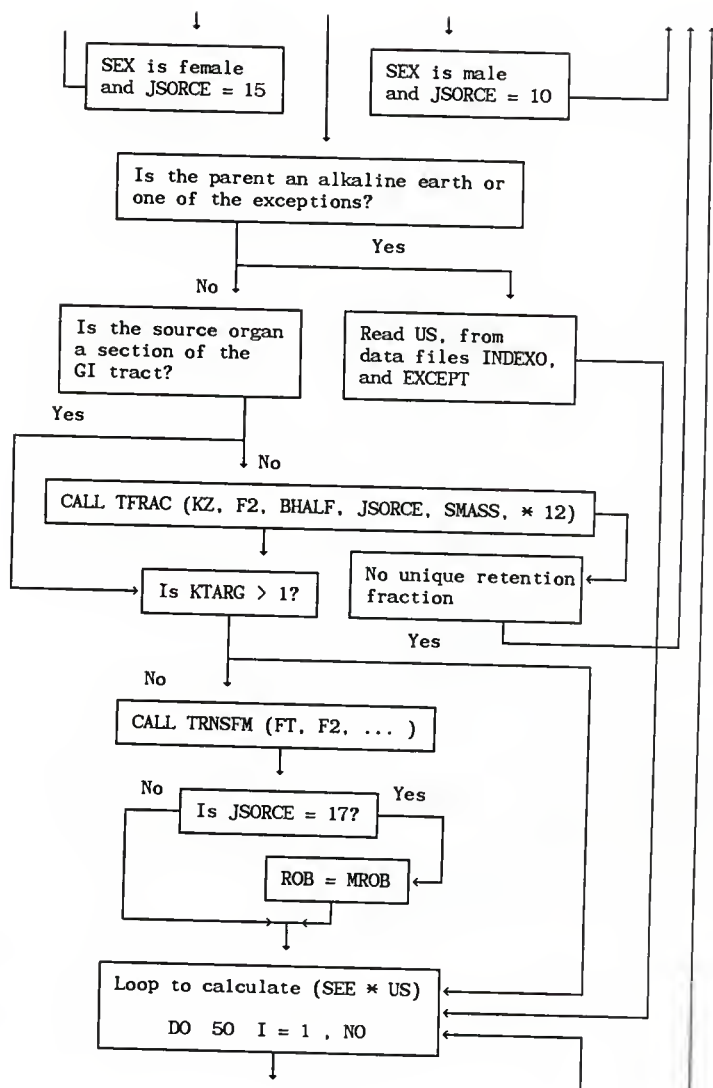
Target Organ Loop
DO 25 KTARG = 1, 19

SEX is female
and KTARG = 3

SEX is male
and KTARG = 4

TMASS = REFMAN (KTARG)

Source Organ Loop
DO 30 JSORCE = 17, 1, -1



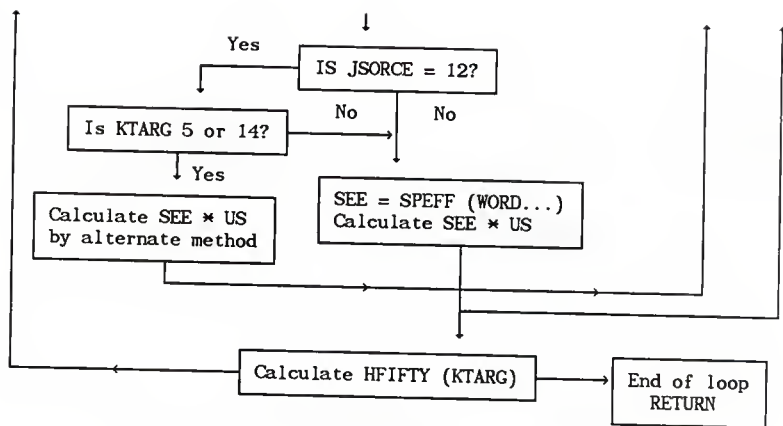


FIG. 3.6. Flow Diagram for "INGEST FORTRAN"

After receiving all the values that were originally requested, the main program now writes these results onto the named files. To begin with, if the source-organ transformations were asked (i.e., MAIS = 1) in the twelfth screen prompt, then the values of US are written to the file FFILE(4). Again, if the mode of data entry was keyboard, then these values are also written on screen for user's convenience.

Before the results of dose commitment can be written onto the named file, a subroutine "RESULT" is called to evaluate weighted committed dose and other quantities as explained below.

3.4.20. RESULT FORTRAN:

The purpose of this subroutine is to evaluate the weighted committed dose equivalent using the 10% exclusion principle. In other words, weighted dose equivalents that are reported are greater than or equal to 10% of the maximum weighted dose. The subroutine also calculates the annual limit on intake. In addition, if the mode of intake is inhalation, it determines the derived air concentration of the nuclide. The arguments of this subroutine are:

- HFIFTY(24) - This vector of maximum length 24, represents the specific committed dose equivalent (Sv/Bq) in 19 target organs or tissues.
- WDOSE(24) - Also a vector of length 24, it describes the weighted committed dose equivalent (Sv/Bq) in 19 target organs or tissues which are greater than or equal to 10% of the maximum.

- ALI - This is a real variable which gives the annual limit on intake (Bq) of the radionuclide. It is equal to variable POST. However, when the annual limit on intake is determined by the nonstochastic limit, this variable stands for the greatest value of the annual intake (Bq) that satisfies the Commission's recommendation for limiting non-stochastic effects.
- POST - This real variable describes the greatest value of annual intake (Bq) that proscribes to the criterion for limiting stochastic effects as recommended by the ICRP-30.
- IRGANT - When the annual limit on intake is determined by the non-stochastic limit on dose equivalent in a particular organ or tissue, that organ or tissue number (according to Table 3.1) is given by this integer variable.
- DAC - This variable gives the concentration of the radionuclide which if inhaled at the (occupational) rate of 9600 liters per day, 5 days per week, 50 weeks per year, would lead to the annual limit of intake for inhalation (Bq/m^3).
- KZ - Atomic number of the nuclide.
- REMDR - Maximum committed dose equivalent in a target organ or tissue which is not included in the metabolic model, GI tract model, and the table of weighting factors shown in Table 1.1.
- WTF - This is a weighting factor determined by the number of target organs and tissues, up to a maximum of 5, which

are not eliminated under the 10% rule or included in metabolic model, GI tract, and Table 1.1, which also qualify for a weighting factor of 0.06.

WREMDR - Weighted committed dose equivalent that is obtained by multiplying REMDR and WTF.

SUM - Sum of all the weighted committed dose equivalents (Sv/Bq).

This subroutine first finds a maximum of target organs or tissues collectively called "remainder" that are not included in organs in Table 1.1, receiving the highest dose equivalents; the exposure of all other remaining tissues is neglected. After that, the weighted committed dose equivalent in each target organ is calculated by multiplying the specific committed dose equivalent with respective weights.

The weighted doses are then compared with the maximum, and if any of the target organ or tissue has a weighted dose less than 10% of the maximum, it is neglected.

The committed dose equivalent assigned to the "Remainder" as a variable REMDR is the maximum committed dose equivalent in any target or tissue which is not included in the GI model, Table 1.1, and the metabolic model. The metabolic model is checked by using the data file "RETENT". A weighting factor WTF is determined by the number of organs, which are not eliminated under the 10% rule or included in the above categories, which also qualify for a weighting factor of 0.06. In the event of no such organ or tissues, no committed dose equivalent,

or weighted committed dose equivalent WREMDR which is merely the product of REMDR and WTF, is given for remainder tissues.

The annual limit on intake for occupational exposure is calculated as the greatest value which satisfies both the stochastic and non-stochastic limit set forth by the Commission in ICRP-30. If the non-stochastic limit overrides the stochastic, then the value that meets both the criteria are reported with the particular organ or tissue concerned.

With the value of ALI, the derived air concentration is then calculated in the case of inhalation, and all these values returned to the main program.

3.4.21 SUBMER FORTRAN:

This is the third possible route from the subroutine "ICRP" when the value of integer variable INTAKE equals 3. This subroutine opens data files "LIST" and "NOBLE" for dose equivalent rate in target organs from submersion in a semi-infinite cloud of radioactive noble gas or elemental tritium.

The arguments are:

- WORD - Identification of the radionuclide.
- HRATE(24) - A vector of length 24, it represents the dose equivalent rate in target organs or tissues from submersion in unit concentration of the isotope.
- DER - A real variable representating derived air concentration (DAC).

- RISK - A real variable that describes derived air concentration when determined by the non-stochastic limit.
- ORGAN - An integer variable which gives the organ number when DAC is determined by consideration of non-stochastic effects.

A search is done in data file "LIST" by comparing the variable WORD with the nuclides in the file. In the event of a proper match, a record number beside the identification of nuclide is retrieved. This record number is the pointer to dose equivalent rates and DAC for that nuclide in data file "EXCEPT". After retrieval of these quantities, the values are returned to the subroutine "ICRP" which directs them to the main program "DOSE".

The results from the above subroutine are then written onto the file FFILE(2) named by the user. After that according to the data input in earlier screens, the program is either stopped automatically, returned to the fourth screen for more data entry or manual stop, or reads more data from the input file.

3.5 RESULTS AND DISCUSSION

Sample calculations of specific effective energy in a target organ from each transformation in a source organ is shown in Table 3.2, 3.3, and 3.4 for radionuclides ^{131}I , $^{113\text{m}}\text{In}$, and ^{121}Te respectively. These radionuclides were chosen due to their different modes of radioactive decay. ^{131}I decays by emission of a beta particle into ^{131}Xe . ^{121}Te decays by positron emission into ^{121}Sb , and $^{113\text{m}}\text{In}$ goes through

isomeric transition from metastable to stable state by emission of gamma radiation.

Results of calculations of specific committed dose equivalents, along with the annual limits of intake, are presented in Tables 3.5 - 3.13 for selected radionuclides. Both modes of intake, ingestion and inhalation are considered in these tables. These results are compared against the published values of the ICRP-30 in Tables 3.14 - 3.17 for a few radionuclides. The ICRP-30 selects data for inclusion in the tables by applying the 10% rule for exclusion of target organs. Only those values are reported which are greater than or equal to 10% of the maximum. Hence, comparison is possible only for selected target organs in tables 3.14 - 3.17.

One can see from the Tables 3.14 - 3.17 that except for three radionuclides, namely, ^{89}Sr , $^{99\text{m}}\text{Tc}$, and $^{131\text{m}}\text{Te}$, all values of specific committed dose equivalents are almost identical (differences in decay schemes used by the two programs account for these minor discrepancies) to the results of ICRP-30. The differences in the exceptions are explained below:

a) ^{89}Sr : To understand the difference in the specific committed dose equivalent of the target organ red marrow, first the S-table for ^{89}Sr was checked against the one in ICRP-30. The results were identical. After that, the table of number of transformations in source organs was compared. It seemed that the difference was due to the number of transformations in the source organ mineral bone. In the case of ingestion for $f_1 = 0.3$, the ICRP-30 reports a value of $1.7\text{E}05$ for the

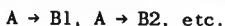
number of transformations per unit intake of activity in cortical bone and a value of 1.4E05 for trabecular bone as source organ. The half life of ^{89}Sr is 50.5 days. Now, according to the criteria described in Part I of Publication 30, isotopes of alkaline earth elements with radioactive half-lives greater than 15 days should be considered to be uniformly distributed throughout the volume of mineral bone, and for radionuclides assumed to be uniformly distributed throughout the volume of mineral bone, $U_{\text{trabecular}} = 0.2 U_{\text{mineral}}$ and $U_{\text{cortical}} = 0.8 U_{\text{mineral}}$. We can see that there is a discrepancy between stated and applied ICRP criteria since the values that are given for U_{cortical} , the number of transformations per unit intake of activity in the cortical bone, and $U_{\text{trabecular}}$ are not in the required ratio. The same is the case with inhalation as the mode of intake.

b) $^{99\text{m}}\text{Tc}$: The difference in the results of specific committed dose equivalents for stomach wall as the target organ, both in the case of ingestion and inhalation, is expected. As described earlier, there is an anomaly in the case of isotopes of elements technetium and rhenium. The ICRP-30 reports both the stomach content and the stomach wall as source organs, and describes transformations for each of them. This is in contrast to all other radionuclides in which only the stomach content is treated as the source organ. In this program, the transformations per unit intake of activity given in the ICRP-30 for both the stomach content and the stomach wall were summed and treated as only one source organ, the stomach content.

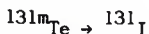
c) ^{131}Te : The difference in this case is due to the mode of radioactive decay. This program, as mentioned earlier, considers only one major branch, i.e.,



whereas, the ICRP program can consider a decay mode like



For the vast majority of nuclides, this lack of feature does not affect the result appreciably since one major mode of decay is usually predominant. But in this special case, ^{131m}Te decays 77.8% of the time into ^{131}I and 22.8% of the time into ^{131}Te . But this program can consider only the major path, i.e.,



and thus will underpredict the results.

d) Features: As explained earlier, the whole program is divided into separate subroutines for calculation of important quantities. The central code "DOSE" merely calls the subroutine "ICRP" with the given input data and prints the output data received from the "ICRP" in a prescribed format. The "ICRP" subroutine, in turn, calls different subroutines for calculation of different quantities according to the given input. This feature of independent subroutines allows more freedom to manipulate or modify the program, e.g., if the specific committed dose equivalent is described in any other subject other than the "reference man", the masses of source and target organs can be changed in the appropriate subroutines.

This convenience of alteration is also provided in the case of biological and radiological decay data. In the case of biological decay, each element is linked with five lines of records in the data files "BFFRAC" and "RETENT" with the unused ones as blanks, so that new data can be entered or the present data can be changed easily. Also with the use of pointers described earlier, an appendage or change in the data files on radiological decay can be done easily.

TABLE 3.2. I-131 S-FACTORS FOR THE ADULT - MeV/g

TARGET ORGAN	SOURCE ORGAN								
	Bladder	Stomach	SI	ULI	LLI	Kidneys	Liver	Lungs	Muscle
Bladder	5.73E-04	4.84E-07	4.02E-06	2.67E-06	7.97E-06	4.75E-07	3.50E-07	8.68E-08	2.36E-06
Stomach	4.20E-07	4.49E-04	4.66E-06	4.85E-06	2.37E-06	4.46E-06	2.54E-06	2.46E-06	1.86E-06
SI	3.61E-06	3.47E-06	2.79E-04	2.19E-05	1.21E-05	3.70E-06	2.15E-06	3.29E-07	2.06E-06
ULI	3.10E-06	4.48E-06	3.07E-05	4.89E-04	5.60E-06	3.82E-06	3.33E-06	4.33E-07	2.16E-06
LLI	9.37E-06	1.70E-06	9.19E-06	4.00E-06	7.73E-04	1.15E-06	3.81E-07	1.23E-07	2.25E-06
Kidneys	4.52E-07	4.49E-06	4.13E-06	3.63E-06	1.18E-06	7.65E-04	5.01E-06	1.30E-06	1.89E-06
Liver	3.40E-07	2.66E-06	2.42E-06	3.35E-06	4.28E-07	5.21E-06	1.39E-04	3.21E-06	1.46E-06
Lungs	5.41E-08	2.37E-06	4.02E-07	4.20E-07	1.31E-07	1.21E-06	3.22E-06	2.10E-04	1.77E-06
Muscle	2.36E-06	1.86E-06	2.06E-06	1.96E-06	2.25E-06	1.89E-06	1.47E-06	1.76E-06	5.92E-06
Ovaries	8.94E-06	6.88E-07	1.27E-05	1.60E-05	2.34E-05	1.60E-06	4.54E-07	1.88E-07	2.64E-06
Pancreas	3.71E-07	2.35E-05	2.73E-06	2.73E-06	9.66E-07	8.63E-06	5.82E-06	3.52E-06	2.38E-06
Bone Surf	8.72E-07	2.24E-06	1.16E-06	1.06E-06	1.52E-06	1.40E-06	1.07E-06	1.42E-06	1.44E-06
Red Marr.	1.91E-06	1.53E-06	3.75E-06	3.23E-06	4.58E-06	3.59E-06	1.56E-06	1.80E-06	1.95E-06
Skin	8.21E-07	7.22E-07	6.68E-07	6.86E-07	7.51E-07	8.38E-07	7.64E-07	8.54E-07	1.14E-06
Spleen	2.65E-07	1.29E-05	2.07E-06	1.73E-06	1.17E-06	1.13E-05	1.30E-06	2.92E-06	1.95E-06
Testes	6.41E-06	6.36E-08	4.86E-07	5.83E-07	2.69E-06	1.83E-07	1.44E-07	2.70E-08	1.61E-06
Thyroid	9.95E-09	1.87E-07	4.49E-08	4.85E-08	1.97E-08	1.12E-07	2.68E-07	1.41E-06	1.80E-06
Uterus	2.01E-05	1.15E-06	1.19E-05	6.07E-06	8.19E-06	1.24E-06	5.55E-07	1.30E-07	2.80E-06
Tot. Body	2.79E-06	3.17E-06	4.80E-06	3.84E-06	4.14E-06	4.96E-06	4.98E-06	4.65E-06	4.60E-06

SOURCE ORGAN

TARGET ORGAN	Ovaries	Pancreas	Trab Bone	Cort Bone	Skin	Spleen	Testes	Thyroid	Tot. Body
Bladder	8.96E-06	2.35E-07	7.61E-07	7.61E-07	7.95E-07	2.12E-07	6.60E-06	1.01E-08	5.11E-06
Stomach	1.08E-06	2.34E-05	7.71E-07	7.71E-07	8.14E-07	1.28E-05	1.16E-07	1.24E-07	5.15E-06
SI	1.57E-05	2.39E-06	1.05E-06	1.05E-06	7.00E-07	1.86E-06	6.47E-07	1.64E-08	5.35E-06
ULI	1.48E-05	2.89E-06	9.58E-07	9.58E-07	7.15E-07	1.77E-06	4.57E-07	1.65E-08	5.23E-06
LLI	1.89E-05	7.32E-07	1.33E-06	1.33E-06	7.50E-07	8.88E-07	3.71E-06	1.64E-08	5.19E-06
Kidneys	1.41E-06	8.31E-06	1.23E-06	1.23E-06	9.43E-07	1.15E-05	1.19E-07	6.43E-08	5.04E-06
Liver	8.02E-07	5.58E-06	9.22E-07	9.22E-07	8.43E-07	1.40E-06	6.61E-08	1.92E-07	4.98E-06
Lungs	1.30E-07	3.23E-06	1.31E-06	1.31E-06	8.83E-07	2.93E-06	1.88E-08	1.36E-06	4.71E-06
Muscle	2.64E-06	2.39E-06	1.44E-06	1.44E-06	1.14E-06	1.95E-06	1.61E-06	1.80E-06	4.60E-06
Ovaries	2.40E-02	5.09E-07	1.23E-06	1.23E-06	5.20E-07	1.12E-06	0.00E-01	1.94E-08	5.03E-06
Pancreas	7.28E-07	3.43E-03	1.30E-06	1.30E-06	7.68E-07	2.54E-05	7.47E-08	1.12E-07	5.24E-06
Bone Surf	1.37E-06	1.34E-06	3.87E-04	3.86E-04	1.14E-06	1.11E-06	9.35E-07	1.05E-06	4.80E-06
Red Marr.	4.65E-06	2.55E-06	6.78E-05	4.29E-06	1.07E-06	1.66E-06	7.53E-07	1.16E-06	5.05E-06
Skin	6.62E-07	6.38E-07	1.08E-06	1.08E-06	6.98E-05	7.74E-07	2.04E-06	1.12E-06	3.86E-06
Spleen	8.43E-07	2.54E-05	1.02E-06	1.02E-06	8.40E-07	1.26E-03	1.11E-07	1.71E-07	5.09E-06
Testes	0.00E-01	9.43E-08	7.95E-07	7.95E-07	1.25E-06	1.13E-07	5.60E-03	3.46E-09	4.79E-06
Thyroid	1.94E-08	2.20E-07	1.34E-06	1.34E-06	1.07E-06	1.80E-07	3.45E-09	1.03E-02	4.53E-06
Uterus	2.55E-05	8.42E-07	8.11E-07	8.11E-07	6.72E-07	5.91E-07	0.00E-01	1.83E-08	5.28E-06
Tot. Body	5.39E-06	5.34E-06	4.63E-06	4.63E-06	3.88E-06	4.97E-06	4.59E-06	4.44E-06	4.62E-06

TABLE 3.3. IN-113M S-FACTORS FOR THE ADULT - MeV/g

TARGET ORGAN	SOURCE ORGAN								
	Bladder	Stomach	SI	ULI	LLI	Kidneys	Liver	Lungs	Muscle
Bladder	3.98E-04	3.30E-07	2.79E-06	1.78E-06	5.40E-06	3.17E-07	2.48E-07	5.82E-08	1.61E-06
Stomach	2.72E-07	3.12E-04	3.15E-06	3.32E-06	1.61E-06	3.01E-06	1.71E-06	1.67E-06	1.28E-06
SI	2.42E-06	2.35E-06	1.94E-04	1.53E-05	8.44E-06	2.49E-06	1.45E-06	2.21E-07	1.40E-06
ULI	2.11E-06	3.05E-06	2.23E-05	3.39E-04	3.92E-06	2.57E-06	2.27E-06	2.93E-07	1.48E-06
LLI	6.39E-06	1.14E-06	6.47E-06	2.74E-06	5.35E-04	7.74E-07	2.60E-07	8.08E-08	1.54E-06
Kidneys	3.08E-07	3.02E-06	2.79E-06	2.43E-06	7.82E-07	5.33E-04	3.40E-06	8.81E-07	1.30E-06
Liver	2.29E-07	1.80E-06	1.63E-06	2.26E-06	2.87E-07	3.55E-06	9.70E-05	2.20E-06	1.00E-06
Lungs	3.54E-08	1.62E-06	2.72E-07	2.81E-07	8.56E-08	8.12E-07	2.18E-06	1.46E-04	1.23E-06
Muscle	1.62E-06	1.27E-06	1.41E-06	1.34E-06	1.53E-06	1.30E-06	1.00E-06	1.23E-06	4.10E-06
Ovaries	6.10E-06	4.19E-07	8.63E-06	1.13E-05	1.63E-05	1.05E-06	2.72E-07	1.26E-07	1.83E-06
Pancreas	2.46E-07	1.62E-05	1.84E-06	1.80E-06	6.50E-07	5.83E-06	3.99E-06	2.41E-06	1.65E-06
Bone Surf	5.81E-07	1.44E-06	7.82E-07	7.14E-07	1.05E-06	9.51E-07	7.25E-07	9.74E-07	9.78E-07
Red Marr.	1.27E-06	1.03E-06	2.54E-06	2.20E-06	3.20E-06	2.44E-06	1.05E-06	1.23E-06	1.36E-06
Skin	5.54E-07	4.90E-07	4.49E-07	4.64E-07	5.08E-07	5.72E-07	5.18E-07	5.80E-07	7.97E-07
Spleen	1.75E-07	8.74E-06	1.40E-06	1.15E-06	7.81E-07	7.78E-06	6.68E-07	1.98E-06	1.34E-06
Testes	4.35E-06	3.52E-08	3.19E-07	3.95E-07	1.84E-06	1.23E-07	9.14E-08	1.79E-08	1.10E-06
Thyroid	6.53E-09	1.25E-07	3.00E-08	3.24E-08	1.31E-08	7.53E-08	1.78E-07	9.49E-07	1.27E-06
Uterus	1.38E-05	7.72E-07	8.12E-06	4.07E-06	5.49E-06	8.27E-07	3.76E-07	8.58E-08	1.91E-06
Tot. Body	1.94E-06	2.20E-06	3.32E-06	2.67E-06	2.87E-06	3.44E-06	3.45E-06	3.23E-06	3.19E-06

SOURCE ORGAN

TARGET ORGAN	Ovaries	Pancreas	Trab Bone	Cort Bone	Skin	Spleen	Testes	Thyroid	Tot. Body
Bladder	6.03E-06	1.62E-07	5.26E-07	5.26E-07	5.33E-07	1.31E-07	4.46E-06	6.61E-09	3.54E-06
Stomach	7.31E-07	1.63E-05	5.22E-07	5.22E-07	5.52E-07	8.78E-06	8.18E-08	8.54E-08	3.57E-06
SI	1.09E-05	1.60E-06	7.07E-07	7.07E-07	4.71E-07	1.25E-06	4.38E-07	9.34E-09	3.70E-06
ULI	1.05E-05	1.94E-06	6.41E-07	6.41E-07	4.81E-07	1.19E-06	3.01E-07	9.60E-09	3.63E-06
LLI	1.34E-05	4.89E-07	8.92E-07	8.92E-07	5.04E-07	6.01E-07	2.53E-06	1.09E-08	3.60E-06
Kidneys	9.55E-07	5.63E-06	8.16E-07	8.16E-07	6.43E-07	7.94E-06	7.91E-08	4.03E-08	3.48E-06
Liver	5.37E-07	3.78E-06	6.22E-07	6.22E-07	5.73E-07	9.38E-07	4.23E-08	1.29E-07	3.45E-06
Lungs	8.66E-08	2.18E-06	8.83E-07	8.83E-07	5.98E-07	2.00E-06	1.21E-08	9.11E-07	3.28E-06
Muscle	1.83E-06	1.65E-06	9.76E-07	9.76E-07	7.95E-07	1.34E-06	1.10E-06	1.27E-06	3.19E-06
Ovaries	1.66E-02	3.13E-07	8.48E-07	8.48E-07	3.30E-07	7.90E-07	0.00E-01	1.28E-08	3.50E-06
Pancreas	4.73E-07	2.38E-03	8.70E-07	8.70E-07	5.22E-07	1.78E-05	4.61E-08	7.47E-08	3.63E-06
Bone Surf	9.27E-07	8.99E-07	3.40E-05	2.32E-05	7.96E-07	7.50E-07	6.24E-07	7.07E-07	3.35E-06
Red Marr.	3.15E-06	1.71E-06	4.69E-05	3.27E-06	7.45E-07	1.12E-06	5.01E-07	7.77E-07	3.52E-06
Skin	4.45E-07	4.28E-07	7.41E-07	7.41E-07	4.82E-05	5.25E-07	1.45E-06	7.71E-07	2.67E-06
Spleen	5.73E-07	1.78E-05	6.94E-07	6.94E-07	5.63E-07	8.81E-04	8.07E-08	1.12E-07	3.52E-06
Testes	0.00E-01	6.02E-08	5.29E-07	5.29E-07	8.69E-07	7.59E-08	3.89E-03	2.22E-09	3.33E-06
Thyroid	1.29E-08	1.56E-07	9.38E-07	9.38E-07	7.42E-07	1.21E-07	2.22E-09	7.17E-03	3.15E-06
Uterus	1.75E-05	5.66E-07	5.29E-07	5.29E-07	4.48E-07	3.91E-07	0.00E-01	1.21E-08	3.61E-06
Tot. Body	3.73E-06	3.69E-06	3.21E-06	3.21E-06	2.68E-06	3.44E-06	3.18E-06	3.08E-06	3.21E-06

TABLE 3.4. TE-121 S-FACTORS FOR THE ADULT - MeV/g

	SOURCE ORGAN								
TARGET ORGAN	Bladder	Stomach	SI	ULI	LLI	Kidneys	Liver	Lungs	Muscle
Bladder	1.83E-04	8.85E-07	6.30E-06	3.73E-06	1.10E-05	7.53E-07	6.30E-07	1.46E-07	3.55E-06
Stomach	6.65E-07	1.30E-04	6.70E-06	7.23E-06	3.46E-06	6.51E-06	3.68E-06	3.66E-06	2.81E-06
SI	5.06E-06	5.09E-06	7.77E-05	3.37E-05	1.84E-05	5.32E-06	3.06E-06	5.41E-07	3.07E-06
ULI	4.46E-06	6.44E-06	4.96E-05	1.13E-04	8.56E-06	5.56E-06	4.98E-06	7.40E-07	3.25E-06
LLI	1.39E-05	2.46E-06	1.41E-05	6.10E-06	1.45E-04	1.68E-06	6.07E-07	1.53E-07	3.36E-06
Kidneys	7.30E-07	6.36E-06	6.14E-06	5.28E-06	1.61E-06	1.95E-04	7.35E-06	2.01E-06	2.90E-06
Liver	5.47E-07	3.92E-06	3.51E-06	4.80E-06	6.68E-07	7.83E-06	6.01E-05	4.76E-06	2.23E-06
Lungs	1.00E-07	3.64E-06	6.48E-07	6.67E-07	2.08E-07	1.85E-06	4.64E-06	4.20E-05	2.75E-06
Muscle	3.55E-06	2.81E-06	3.08E-06	2.94E-06	3.36E-06	2.90E-06	2.24E-06	2.75E-06	3.32E-06
Ovaries	1.19E-05	1.19E-06	1.81E-05	2.44E-05	3.37E-05	2.31E-06	4.93E-07	2.97E-07	4.03E-06
Pancreas	5.04E-07	3.54E-05	3.98E-06	3.49E-06	1.44E-06	1.29E-05	9.28E-06	5.32E-06	3.65E-06
Bone Surf	1.13E-06	1.20E-06	1.57E-06	1.44E-06	2.20E-06	1.99E-06	1.49E-06	2.07E-06	2.17E-06
Red Marr.	2.42E-06	2.04E-06	5.08E-06	4.40E-06	6.76E-06	5.06E-06	2.13E-06	2.59E-06	2.89E-06
Skin	1.28E-06	1.16E-06	1.04E-06	1.08E-06	1.15E-06	1.31E-06	1.18E-06	1.33E-06	1.87E-06
Spleen	4.40E-07	1.88E-05	3.02E-06	2.39E-06	1.83E-06	1.69E-05	1.95E-06	4.27E-06	2.98E-06
Testes	9.84E-06	7.87E-08	7.29E-07	1.00E-06	3.88E-06	2.92E-07	2.51E-07	4.95E-08	2.46E-06
Thyroid	1.98E-08	2.96E-07	7.99E-08	8.55E-08	3.70E-08	1.86E-07	3.55E-07	2.35E-06	2.87E-06
Uterus	3.00E-05	1.72E-06	1.74E-05	8.72E-06	1.09E-05	1.60E-06	8.31E-07	1.88E-07	4.08E-06
Tot. Body	3.84E-06	3.68E-06	4.10E-06	3.96E-06	3.96E-06	3.60E-06	3.63E-06	3.14E-06	3.08E-06

SOURCE ORGAN

TARGET ORGAN	Ovaries	Pencreas	Treb Bone	Cort Bone	Skin	Spleen	Testes	Thyroid	Tot. Body
Bladder	1.31E-05	3.61E-07	1.21E-06	1.21E-06	1.27E-06	3.07E-07	9.99E-06	2.00E-08	3.89E-06
Stomach	1.59E-06	3.51E-05	1.12E-06	1.12E-06	1.28E-06	1.94E-05	2.17E-07	2.48E-07	3.74E-06
SI	2.40E-05	3.39E-06	1.57E-06	1.57E-06	1.08E-06	2.66E-06	1.02E-06	2.16E-08	4.15E-06
ULI	2.31E-05	4.15E-06	1.37E-06	1.37E-06	1.15E-06	2.70E-06	6.34E-07	2.81E-08	4.10E-06
LLI	2.97E-05	1.03E-06	1.89E-06	1.89E-06	1.18E-06	1.33E-06	5.39E-06	3.10E-08	4.14E-06
Kidneys	2.16E-06	1.21E-05	1.83E-06	1.83E-06	1.49E-06	1.72E-05	2.06E-07	1.11E-07	3.71E-06
Liver	1.18E-06	8.02E-06	1.38E-06	1.38E-06	1.33E-06	2.05E-06	1.11E-07	3.13E-07	3.70E-06
Lungs	2.26E-07	4.65E-06	1.93E-06	1.93E-06	1.35E-06	4.37E-06	3.11E-08	2.07E-06	3.31E-06
Muscle	4.03E-06	3.65E-06	2.17E-06	2.17E-06	1.86E-06	2.97E-06	2.46E-06	2.86E-06	3.09E-06
Ovaries	2.81E-03	8.67E-07	2.11E-06	2.11E-06	6.54E-07	1.92E-06	0.00E-01	3.69E-08	3.52E-06
Pencreas	1.15E-06	6.28E-04	1.77E-06	1.77E-06	1.11E-06	4.01E-05	8.74E-08	1.28E-07	3.67E-06
Bone Surf	1.87E-06	1.82E-06	1.99E-05	1.97E-05	1.79E-06	1.56E-06	1.28E-06	1.48E-06	3.36E-06
Red Marr.	6.32E-06	3.42E-06	1.02E-05	7.49E-06	1.66E-06	2.25E-06	1.01E-06	1.63E-06	3.69E-06
Skin	1.02E-06	9.87E-07	1.70E-06	1.70E-06	7.44E-06	1.24E-06	3.32E-06	1.76E-06	1.96E-06
Spleen	1.31E-06	3.96E-05	1.65E-06	1.65E-06	1.32E-06	3.43E-04	2.36E-07	2.64E-07	3.77E-06
Testes	0.00E-01	1.34E-07	1.13E-06	1.13E-06	1.79E-06	1.87E-07	1.05E-03	7.39E-09	3.86E-06
Thyroid	3.69E-08	3.22E-07	2.02E-06	2.02E-06	1.66E-06	2.88E-07	7.38E-09	1.55E-03	3.11E-06
Uterus	3.62E-05	1.20E-06	1.20E-06	1.20E-06	1.14E-06	9.29E-07	0.00E-01	3.50E-08	3.84E-06
Tot. Body	4.20E-06	4.12E-06	3.12E-06	3.12E-06	2.01E-06	3.61E-06	3.05E-06	2.85E-06	3.11E-06

TABLE 3.5 Specific committed dose equivalents (Sv/Bq) for selected ingested radionuclides and their daughters.
Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks.

Nuclide	Na-24	P-32	K-40	Cr-51	Cr-51	Mn-54	Mn-56	Fe-55	Fe-59	Co-58	Co-60
ALI (Bq)	1.0	0.8	1.0	0.1	0.01	0.1	0.1	0.1	0.1	0.05	0.05
ALI (Bq)	1.3E+08	2.4E+07	1.1E+07	1.4E+09	1.3E+09	6.8E+07	2.0E+08	3.2E+08	2.8E+07	6.4E+07	1.8E+07
Lungs	2.6E+10	6.6E+10	4.3E+09	4.3E+12	9.5E+13	2.3E+10	8.8E+12	1.0E+10	6.4E+10	8.5E+11	8.8E+10
Thyroid	2.6E+10	6.6E+10	4.3E+09	3.6E+12	4.2E+13	1.3E+10	2.4E+12	1.0E+10	6.1E+10	6.3E+11	7.9E+10
Testes	0.0E+01	0.0E+01	0.0E+01	0.0E+01	0.0E+01	0.0E+01	0.0E+01	0.0E+01	0.0E+01	0.0E+01	0.0E+01
Ovaries	3.4E+10	6.6E+10	4.5E+09	4.0E+11	3.9E+11	9.5E+10	8.5E+11	1.0E+10	1.7E+09	1.0E+09	3.2E+09
Red Marrow	3.7E+10	7.8E+09	4.3E+09	1.3E+11	8.7E+12	5.1E+10	2.4E+11	1.0E+10	8.5E+10	2.6E+10	1.3E+09
Stomach wall	1.2E+09	1.5E+09	4.8E+09	1.9E+11	1.6E+11	4.1E+10	9.0E+10	1.1E+10	1.1E+09	3.9E+10	1.6E+09
SI + contents	3.2E+10	1.1E+09	4.4E+09	4.8E+11	4.6E+11	9.9E+10	1.1E+09	1.1E+10	2.1E+09	1.1E+09	3.6E+09
ULI wall	3.1E+10	3.0E+09	4.4E+09	1.2E+10	1.3E+10	1.4E+09	1.4E+09	1.9E+10	4.0E+09	2.0E+09	5.8E+09
LLI wall	3.4E+10	7.2E+09	4.4E+09	2.8E+10	3.1E+10	2.2E+09	5.3E+10	3.6E+10	8.4E+09	4.0E+09	1.1E+08
Liver	2.9E+10	6.6E+10	4.4E+09	7.0E+12	3.6E+12	1.0E+09	2.6E+11	3.3E+10	1.5E+09	2.5E+10	2.3E+09
Kidneys	3.0E+10	6.6E+10	4.4E+09	8.5E+12	5.0E+12	3.8E+10	3.2E+11	1.0E+10	9.1E+09	2.1E+10	1.4E+09
Bladder wall	3.0E+10	6.6E+10	4.4E+09	1.5E+11	1.2E+11	3.7E+10	2.6E+11	1.0E+10	1.1E+09	3.7E+10	1.8E+09
Muscle	2.7E+10	6.6E+10	4.3E+09	7.5E+12	4.5E+12	2.6E+10	1.8E+11	1.0E+10	7.4E+10	1.8E+10	1.1E+09
Bone Surface	4.7E+10	7.8E+09	4.3E+09	7.9E+12	3.3E+12	5.6E+10	1.1E+11	1.0E+10	6.6E+10	1.3E+11	9.4E+10
Skin	2.1E+10	6.6E+10	4.2E+09	3.9E+12	1.6E+12	1.6E+10	7.8E+12	1.0E+10	5.0E+10	8.5E+10	6.9E+10
Spleen	3.1E+10	6.6E+10	4.3E+09	7.4E+12	3.6E+12	2.7E+10	3.5E+11	5.5E+10	1.8E+09	1.7E+10	1.2E+09
Uterus	3.3E+10	6.6E+10	4.4E+09	1.9E+11	1.6E+11	5.0E+10	5.9E+11	1.0E+10	1.3E+09	4.8E+10	2.1E+09
Pancreas	4.4E+10	6.6E+10	4.4E+09	8.9E+12	5.0E+12	3.8E+10	5.6E+11	1.0E+10	9.1E+09	2.0E+10	1.3E+09
Total Body	3.0E+10	1.3E+09	4.3E+09	9.2E+12	6.2E+12	3.4E+10	3.6E+11	1.1E+10	8.0E+10	2.1E+10	1.2E+09
Remainder	4.4E+10		4.4E+09			5.0E+10			4.8E+10	2.1E+09	

Nuclide	Ni-63	Ni-65	Cu-64	Zn-65	Zn-69	Br-83	Br-84	Rb-88	Rb-89	Sr-89	Sr-90
ALI (Bq)	0.05	0.05	0.5	0.5	0.5	1.0	1.0	1.0	1.0	0.3	0.3
ALI (Bq)	3.4E+08	3.0E+08	4.3E+08	1.3E+07	2.2E+09	1.7E+09	7.2E+08	6.8E+08	1.4E+09	2.4E+07	1.3E+06
Lungs	8.5E+11	2.8E+12	1.3E+11	3.1E+09	4.0E+13	7.1E+12	7.1E+12	2.9E+12	3.8E+12	2.3E+10	0.0E+01
Thyroid	8.5E+11	6.8E+13	1.1E+11	3.2E+09	4.0E+13	7.1E+12	5.3E+12	2.4E+12	2.3E+12	2.3E+10	0.0E+01
Testes	0.0E+01	0.0E+01	0.0E+01	0.0E+01	0.0E+01	0.0E+01	0.0E+01	0.0E+01	0.0E+01	0.0E+01	0.0E+01
Ovaries	8.5E+11	2.4E+11	4.8E+11	3.5E+09	4.0E+13	7.1E+12	6.8E+12	2.8E+12	3.4E+12	2.3E+10	0.0E+01
Red Marrow	8.5E+11	7.3E+12	1.9E+11	4.9E+09	4.9E+13	7.1E+12	6.3E+12	2.8E+12	4.4E+12	1.6E+09	1.3E+07
Stomach wall	1.0E+10	6.2E+10	1.7E+10	3.4E+09	2.1E+10	2.9E+10	6.9E+10	7.3E+10	3.7E+10	2.3E+10	0.0E+01
SI + contents	1.3E+10	7.3E+10	2.1E+10	4.3E+09	1.1E+10	7.1E+12	8.7E+12	3.3E+12	4.7E+12	2.3E+10	0.0E+01
ULI wall	3.7E+10	9.4E+10	6.2E+10	4.3E+09	5.9E+11	7.1E+12	9.2E+12	3.5E+12	5.3E+12	7.2E+09	4.2E+09
LLI wall	9.2E+10	3.6E+10	7.5E+10	5.0E+09	9.5E+12	7.1E+12	7.2E+12	2.9E+12	3.6E+12	2.1E+08	1.8E+08
Liver	8.5E+11	7.3E+12	3.7E+11	3.7E+09	4.0E+13	7.1E+12	7.8E+12	3.1E+12	3.1E+12	2.3E+10	0.0E+01
Kidneys	8.5E+11	1.0E+11	2.0E+11	3.9E+09	4.0E+13	7.1E+12	9.2E+12	3.4E+12	5.3E+12	2.3E+10	0.0E+01
Bladder wall	8.5E+11	8.5E+12	2.2E+11	4.1E+09	4.0E+13	7.1E+12	5.9E+12	2.6E+12	2.7E+12	2.3E+10	0.0E+01
Muscle	8.5E+11	5.6E+12	1.6E+11	3.3E+09	4.0E+13	7.1E+12	6.7E+12	2.8E+12	3.4E+12	2.3E+10	0.0E+01
Bone Surface	8.5E+11	2.9E+12	1.4E+11	4.5E+09	4.9E+13	7.1E+12	5.6E+12	2.8E+12	4.6E+12	4.3E+09	4.0E+07
Skin	8.5E+11	2.5E+12	1.1E+11	2.3E+09	4.0E+13	7.1E+12	5.3E+12	2.5E+12	2.5E+12	2.3E+10	0.0E+01
Spleen	8.5E+11	1.2E+11	1.9E+11	3.6E+09	4.0E+13	7.2E+12	1.6E+11	5.3E+12	1.1E+11	2.3E+10	0.0E+01
Uterus	8.5E+11	1.9E+11	2.8E+11	4.7E+09	4.0E+13	7.1E+12	7.0E+12	2.8E+12	3.3E+12	2.3E+10	0.0E+01
Pancreas	8.5E+11	1.8E+11	5.4E+11	3.6E+09	4.0E+13	7.3E+12	2.5E+11	7.6E+12	1.8E+11	2.3E+10	0.0E+01
Total Body	8.8E+11	1.8E+11	2.2E+11	3.4E+09	2.0E+12	7.7E+12	8.4E+12	4.5E+12	4.6E+12	7.1E+10	4.0E+08
Remainder				4.7E+09							4.0E+08

TABLE 3.6 Specific committed dose equivalents (Sv/Bq) for selected ingested radionuclides and their daughters.
Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks.

Nuclide	Y-90	Y-91m	Y-91	Y-92	Y-93	Zr-95	Zr-97	Nb-95	Mo-99	Tc-99m	Tc-101
f1	0.0001	0.0001	0.0001	0.0001	0.0001	0.002	0.002	0.01	0.8	0.8	0.8
ALI (Bq)	1.6E+07	4.8E+09	1.7E+07	9.9E+07	4.1E+07	5.4E+07	2.3E+07	8.2E+07	6.0E+07	3.5E+09	3.3E+09
Lungs	1.2E-14	1.3E-12	1.6E-13	1.4E-12	8.7E-13	2.3E-11	1.8E-11	2.8E-11	2.0E-10	2.9E-12	3.3E-13
Thyroid	1.2E-14	1.2E-13	8.9E-14	1.8E-13	1.3E-13	8.2E-12	2.7E-12	1.2E-11	1.7E-10	8.4E-11	2.6E-14
Testes	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01
Ovaries	1.2E-14	6.9E-12	3.5E-12	2.0E-11	2.2E-11	8.2E-10	6.3E-10	8.1E-10	2.3E-10	9.5E-12	5.1E-13
Red Marrow	3.6E-13	2.2E-12	4.4E-12	4.9E-12	4.9E-12	2.1E-10	1.3E-10	2.0E-10	4.8E-10	6.0E-12	3.4E-13
Stomach wall	1.1E-09	4.9E-11	6.9E-10	1.4E-09	1.3E-09	3.6E-10	1.2E-09	2.8E-10	6.7E-10	3.9E-11	1.5E-10
SI + contents	2.6E-09	3.1E-11	1.7E-09	2.0E-09	2.5E-09	1.1E-09	3.4E-09	9.2E-10	4.7E-10	2.2E-11	2.3E-11
ULI wall	1.4E-08	3.1E-11	1.0E-08	3.3E-09	7.9E-09	3.1E-09	1.2E-08	1.9E-09	1.5E-09	3.6E-11	1.6E-12
LLI wall	3.1E-08	2.4E-11	3.0E-08	1.7E-09	8.7E-09	7.8E-09	1.8E-08	4.0E-09	3.2E-09	2.4E-11	5.3E-13
Liver	3.5E-13	2.5E-12	4.1E-12	4.6E-12	3.6E-12	7.9E-11	8.1E-11	8.4E-11	2.7E-09	5.7E-12	4.4E-13
Kidneys	1.2E-14	3.9E-12	5.0E-13	6.5E-12	4.8E-12	1.1E-10	1.1E-10	1.4E-10	2.7E-09	4.9E-12	7.6E-13
Bladder wall	1.2E-14	2.3E-12	1.2E-12	5.6E-12	6.5E-12	2.4E-10	1.8E-10	2.4E-10	1.9E-10	4.3E-12	1.9E-13
Muscle	1.2E-14	1.8E-12	5.2E-13	3.5E-12	3.1E-12	1.1E-10	8.2E-11	1.1E-10	1.9E-10	3.3E-12	3.2E-13
Bone Surface	3.6E-13	8.7E-13	4.0E-12	1.8E-12	1.8E-12	4.8E-10	4.6E-11	2.9E-10	7.6E-10	4.1E-12	4.2E-13
Skin	1.2E-14	7.1E-13	2.5E-13	1.4E-12	1.2E-12	4.2E-11	3.1E-11	4.4E-11	1.6E-10	1.8E-12	1.2E-13
Spleen	1.2E-14	6.4E-12	4.7E-13	6.5E-12	3.9E-12	8.9E-11	8.1E-11	1.1E-10	2.0E-10	6.2E-12	1.8E-12
Uterus	1.2E-14	5.8E-12	1.6E-12	1.3E-11	1.1E-11	3.3E-10	2.9E-10	3.4E-10	2.1E-10	6.9E-12	5.5E-13
Pancreas	1.2E-14	1.1E-11	5.3E-13	1.0E-11	5.7E-12	1.1E-10	1.1E-10	1.1E-10	2.4E-10	9.3E-12	3.3E-12
Total Body	1.4E-10	2.5E-12	1.2E-10	3.9E-11	7.3E-11	1.5E-10	2.0E-10	1.4E-10	3.0E-10	4.0E-12	9.1E-13
Remainder		1.1E-11								9.3E-12	

Nuclide	Ru-103	Ru-105	Ru-106	Ag-110m	Te-125m	Ta-127	Te-129m	Te-129	Te-131m	Te-131	Te-132
f1	0.05	0.05	0.05	0.05	0.2	0.2	0.2	0.2	0.2	0.2	0.2
ALI (Bq)	6.9E+07	1.9E+08	7.1E+06	1.7E+07	3.5E+07	2.7E+08	1.9E+07	9.7E+08	1.2E+07	1.2E+08	8.4E+06
Lungs	7.4E-11	6.1E-12	1.4E-09	8.3E-10	4.8E-11	2.9E-12	1.6E-10	4.9E-13	4.4E-11	3.3E-12	3.1E-10
Thyroid	6.2E-11	1.7E-12	1.4E-09	1.8E-10	4.3E-11	2.9E-12	1.6E-10	3.4E-13	4.3E-08	4.3E-09	6.0E-08
Testes	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01
Ovaries	5.9E-10	9.5E-11	1.6E-09	3.0E-09	1.3E-10	4.0E-12	2.4E-10	1.6E-12	6.7E-10	1.5E-11	4.5E-10
Red Marrow	1.7E-10	2.3E-11	1.5E-09	9.4E-10	1.4E-09	6.4E-12	3.4E-09	7.5E-13	1.7E-10	6.1E-12	3.5E-10
Stomach wall	3.1E-10	5.0E-10	3.1E-09	1.5E-09	2.1E-10	2.4E-10	6.3E-10	4.0E-10	1.6E-10	6.3E-10	3.3E-10
SI + contents	8.6E-10	7.9E-10	5.5E-09	3.5E-09	4.2E-10	3.9E-10	1.5E-09	2.7E-10	1.2E-09	5.6E-10	4.1E-10
ULI wall	2.5E-09	1.6E-09	2.5E-08	5.9E-09	1.9E-09	1.2E-09	8.6E-09	1.9E-10	2.9E-09	5.4E-10	3.7E-10
LLI wall	6.3E-09	1.2E-09	7.1E-08	1.1E-08	5.4E-09	1.2E-09	2.5E-08	3.6E-11	4.8E-09	9.3E-12	3.6E-09
Liver	1.1E-10	1.9E-11	1.4E-09	8.5E-09	4.6E-11	3.0E-12	1.6E-10	6.7E-13	1.0E-10	5.2E-12	3.3E-10
Kidneys	1.3E-10	2.7E-11	1.5E-09	1.5E-09	4.6E-11	3.1E-12	1.7E-10	8.7E-13	1.3E-10	7.2E-12	3.3E-10
Bladder wall	2.3E-10	2.7E-11	1.5E-09	1.0E-09	5.2E-11	3.2E-12	1.7E-10	6.6E-13	2.2E-10	4.8E-12	3.9E-10
Muscle	1.2E-10	1.6E-11	1.4E-09	7.5E-10	5.0E-11	3.0E-12	1.7E-10	6.0E-13	1.1E-10	4.8E-12	3.2E-10
Bone Surface	9.8E-11	9.0E-12	1.4E-09	4.9E-10	1.4E-08	6.4E-12	7.7E-09	5.6E-13	7.4E-11	3.5E-12	3.1E-10
Skin	6.9E-11	6.4E-12	1.4E-09	3.7E-10	4.2E-11	2.9E-12	1.6E-10	4.1E-13	5.3E-11	2.6E-12	2.4E-10
Spleen	1.2E-10	2.5E-11	1.5E-09	7.1E-10	4.7E-11	3.1E-12	1.6E-10	1.2E-12	9.0E-11	9.1E-12	3.6E-10
Uterus	2.7E-10	5.5E-11	1.5E-09	1.4E-09	5.7E-11	3.4E-12	1.8E-10	1.2E-12	3.2E-10	1.1E-11	4.0E-10
Pancreas	1.3E-10	3.8E-11	1.5E-09	1.6E-09	4.8E-11	3.2E-12	1.6E-10	1.9E-12	1.0E-10	1.4E-11	3.2E-10
Total Body	1.5E-10	3.2E-11	1.7E-09	1.0E-09	1.9E-10	1.4E-11	5.4E-10	4.0E-12	1.6E-10	1.4E-11	3.4E-10
Remainder				1.6E-09							

TABLE 3.7 Specific committed dose equivalents (Sv/Bq) for selected ingested radionuclides and their daughters.
Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks.

Nuclide	Np-239
AI	0.01
ALI (Bq)	6.2E+07

Lungs	2.4E-12
Thyroid	2.8E-13
Testes	0.0E-01
Ovaries	1.6E-10
Red Marrow	5.6E-11
Stomach wall	3.3E-10
SI + contents	8.2E-10
ULI wall	3.7E-09
LLI wall	8.1E-09
Liver	5.4E-11
Kidneys	2.1E-11
Bladder wall	4.7E-11
Muscle	1.7E-11
Bone Surface	1.5E-10
Skin	5.1E-12
Spleen	1.5E-11
Uterus	6.8E-11
Pancreas	2.2E-11
Total Body	5.9E-11
Remainder	

TABLE 3.8 Specific committed dose equivalent (Sv/Bq) for selected inhaled radionuclides and their daughters.
Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks.

Nuclide	Na-24	P-32	P-32	K-40	Cr-51	Cr-51	Cr-51	Mn-54	Mn-54	Mn-56	Mn-56
f1	1.0	0.8	0.8	1.0	0.01	0.01	0.01	0.1	0.1	0.1	0.1
Class	0	W	0	0	0	W	Y	0	W	0	0
ALI (Bq)	1.9E+08	1.4E+07	3.5E+07	1.7E+07	1.7E+09	8.7E+08	7.0E+08	3.6E+07	3.2E+07	5.7E+08	7.8E+08
DAC (Bq/m ³)	7.9E+04	5.9E+03	1.4E+04	7.1E+03	7.0E+05	3.6E+05	2.9E+05	1.5E+04	1.3E+04	2.4E+05	3.2E+05
Lungs	1.2E-09	2.6E-08	2.5E-09	4.1E-09	3.7E-11	3.7E-10	5.2E-10	1.2E-09	6.6E-09	4.4E-10	5.4E-10
Thyroid	1.5E-10	3.4E-10	4.8E-10	2.7E-09	1.7E-11	9.4E-12	9.3E-12	6.5E-10	7.4E-10	1.2E-11	6.2E-12
Testes	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01
Ovaries	1.8E-10	3.4E-10	4.8E-10	2.8E-09	2.7E-11	2.2E-11	2.0E-11	8.9E-10	7.1E-10	2.2E-11	9.4E-12
Red Marrow	2.1E-10	4.0E-09	5.8E-09	2.7E-09	3.1E-11	1.8E-11	1.7E-11	1.8E-09	1.1E-09	2.3E-11	1.0E-11
Stomach wall	3.1E-10	6.7E-10	6.1E-10	2.8E-09	2.4E-11	2.2E-11	2.4E-11	9.4E-10	1.2E-09	1.4E-10	7.6E-11
SI + contents	1.7E-10	5.0E-10	5.5E-10	2.8E-09	3.0E-11	2.7E-11	2.5E-11	1.1E-09	8.4E-10	1.6E-10	8.2E-11
ULI wall	1.6E-10	1.3E-09	8.5E-10	2.6E-09	4.1E-11	6.1E-11	6.3E-11	1.3E-09	1.1E-09	2.0E-10	1.1E-10
LLI wall	1.7E-10	3.1E-09	1.5E-09	2.6E-09	6.6E-11	1.4E-10	1.5E-10	1.2E-09	1.3E-09	8.4E-11	4.3E-11
Liver	1.9E-10	3.4E-10	4.8E-10	2.6E-09	2.1E-11	2.1E-11	2.4E-11	4.6E-09	2.5E-09	4.0E-11	1.7E-11
Kidneys	1.6E-10	3.4E-10	4.8E-10	2.7E-09	2.1E-11	1.2E-11	1.1E-11	1.4E-09	8.9E-10	1.7E-11	8.0E-12
Bladder wall	1.6E-10	3.4E-10	4.6E-10	2.8E-09	2.2E-11	9.2E-12	6.4E-12	7.2E-10	3.8E-10	1.3E-11	4.7E-12
Muscle	1.6E-10	3.4E-10	4.8E-10	2.7E-09	1.9E-11	1.3E-11	1.4E-11	9.1E-10	8.6E-10	1.5E-11	7.8E-12
Bone Surface	2.6E-10	4.0E-09	5.8E-09	2.7E-09	2.6E-11	1.3E-11	1.2E-11	2.5E-09	1.2E-09	2.0E-11	8.2E-12
Skin	1.2E-10	3.4E-10	4.8E-10	2.6E-09	1.3E-11	7.0E-12	6.6E-12	6.2E-10	4.9E-10	1.1E-11	4.8E-12
Spleen	1.8E-10	3.4E-10	4.8E-10	2.7E-09	2.2E-11	1.9E-11	2.2E-11	9.2E-10	1.2E-09	2.0E-11	1.2E-11
Uterus	1.7E-10	3.4E-10	4.8E-10	2.6E-09	2.4E-11	1.2E-11	8.6E-12	9.2E-10	4.9E-10	1.9E-11	7.6E-12
Pancreas	2.3E-10	3.4E-10	4.6E-10	2.7E-09	2.4E-11	2.3E-11	2.6E-11	1.4E-09	1.4E-09	2.4E-11	1.4E-11
Total Body	1.9E-10	1.0E-09	9.7E-10	2.7E-09	2.0E-11	1.9E-11	2.2E-11	1.1E-09	9.9E-10	2.4E-11	1.7E-11
Remainder				2.8E-09	2.4E-11			1.4E-09	1.4E-09		
Nuclide	Fe-55	Fe-55	Fe-59	Fe-59	Co-58	Co-58	Co-60	Co-60	Ni-63	Ni-63	Ni-65
f1	0.1	0.1	0.1	0.1	0.05	0.05	0.05	0.05	0.05	0.05	0.05
Class	0	W	0	W	W	Y	W	Y	W	0	W
ALI (Bq)	7.5E+07	1.4E+08	1.3E+07	1.9E+07	4.1E+07	2.6E+07	6.7E+06	1.2E+06	9.8E+07	5.9E+07	1.1E+09
DAC (Bq/m ³)	3.1E+04	6.3E+04	5.2E+03	7.9E+03	1.7E+04	1.1E+04	2.8E+03	5.0E+02	4.1E+04	2.5E+04	4.6E+05
Lungs	5.1E-10	1.1E-09	3.5E-09	1.4E-08	7.9E-09	1.6E-08	3.6E-08	3.4E-07	3.1E-09	8.8E-10	3.8E-10
Thyroid	4.9E-10	1.7E-10	3.0E-09	1.2E-09	5.5E-10	8.7E-10	3.7E-09	1.6E-08	2.5E-10	8.2E-10	2.3E-12
Testes	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01
Ovaries	4.9E-10	1.7E-10	3.3E-09	1.4E-09	6.5E-10	6.2E-10	4.0E-09	4.8E-09	2.5E-10	8.2E-10	3.3E-12
Red Marrow	4.9E-10	1.7E-10	3.2E-09	1.3E-09	6.3E-10	9.2E-10	4.2E-09	1.7E-08	2.5E-10	8.2E-10	3.0E-12
Stomach wall	4.9E-10	1.7E-10	3.4E-09	1.7E-09	9.0E-10	1.4E-09	5.4E-09	2.7E-08	2.6E-10	8.3E-10	4.8E-11
SI + contents	4.9E-10	1.7E-10	3.9E-09	1.7E-09	7.8E-10	7.5E-10	4.7E-09	7.1E-09	2.7E-10	8.3E-10	5.5E-11
ULI wall	5.0E-10	2.1E-10	4.1E-09	2.6E-09	1.2E-09	1.2E-09	5.9E-09	9.7E-09	3.9E-10	8.7E-10	7.1E-11
LLI wall	5.3E-10	3.0E-10	4.6E-09	4.5E-09	2.0E-09	2.0E-09	8.1E-09	7.9E-09	6.7E-10	9.5E-10	2.8E-11
Liver	1.6E-09	5.5E-10	7.1E-09	2.7E-09	1.1E-09	1.6E-09	9.2E-09	3.4E-08	2.5E-10	8.2E-10	4.1E-12
Kidneys	4.9E-10	1.7E-10	3.8E-09	1.4E-09	5.7E-10	7.6E-10	4.5E-09	1.6E-08	2.5E-10	8.2E-10	3.1E-12
Bladder wall	4.9E-10	1.7E-10	3.7E-09	1.2E-09	3.4E-10	2.4E-10	3.4E-09	3.0E-09	2.5E-10	8.2E-10	2.0E-12
Muscle	4.9E-10	1.7E-10	3.0E-09	1.3E-09	6.1E-10	9.4E-10	4.2E-09	1.8E-08	2.5E-10	8.2E-10	2.4E-12
Bone Surface	4.9E-10	1.7E-10	2.9E-09	1.1E-09	4.8E-10	6.9E-10	3.5E-09	1.4E-08	2.5E-10	8.2E-10	2.0E-12
Skin	4.9E-10	1.7E-10	2.2E-09	8.5E-10	3.3E-10	4.8E-10	2.6E-09	1.0E-08	2.5E-10	8.2E-10	2.0E-12
Spleen	2.7E-09	9.3E-10	8.5E-09	3.0E-09	8.9E-10	1.5E-09	5.2E-09	2.7E-08	2.5E-10	8.2E-10	4.0E-12
Uterus	4.9E-10	1.7E-10	4.0E-09	1.4E-09	4.2E-10	3.1E-10	3.9E-09	4.6E-09	2.5E-10	8.2E-10	3.0E-12
Pancreas	4.9E-10	1.7E-10	3.6E-09	1.8E-09	1.0E-09	1.7E-09	6.0E-09	3.2E-08	2.5E-10	8.2E-10	4.9E-12
Total Body	5.2E-10	1.9E-10	3.1E-09	1.5E-09	7.1E-10	1.1E-09	4.6E-09	2.2E-08	2.9E-10	8.3E-10	9.2E-12
Remainder	5.2E-10		4.0E-09								

TABLE 3.9 Specific committed dose equivalents (Sv/Bq) for alected inhaled radionuclides and their daughters.
Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks.

Nuclide	Ni-65	Cu-64	Cu-64	Cu-64	Zn-65	Zn-69	Br-83	Br-83	Br-84	Br-84	Rb-88
f1	0.05	0.5	0.5	0.5	0.5	0.5	1.0	1.0	1.0	1.0	1.0
Class	0	0	W	Y	Y	Y	0	W	0	W	0
ALI (Bq)	8.7E+08	1.1E+09	8.7E+08	8.0E+08	1.2E+07	5.3E+09	2.5E+09	2.3E+09	2.1E+09	2.4E+09	2.3E+09
DAC (Bq/m ³)	3.6E+05	4.4E+05	3.6E+05	3.3E+05	4.9E+03	2.2E+06	1.0E+06	9.8E+05	8.8E+05	1.0E+06	9.7E+05
Lungs	3.1E-10	2.0E-10	3.4E-10	3.5E-10	2.1E-08	7.8E-11	1.5E-10	1.8E-10	1.6E-10	1.7E-10	1.5E-10
Thyroid	5.5E-12	1.1E-11	6.0E-12	4.9E-12	3.0E-09	2.6E-14	3.2E-12	1.1E-12	3.1E-12	1.4E-12	1.4E-12
Testes	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01
Ovaries	8.5E-12	1.6E-11	1.2E-11	1.2E-11	2.0E-09	2.6E-14	3.2E-12	1.1E-12	2.9E-12	8.6E-13	1.3E-12
Red Marrow	6.7E-12	1.3E-11	7.9E-12	7.0E-12	3.9E-09	3.2E-14	3.2E-12	1.1E-12	3.3E-12	1.5E-12	1.5E-12
Stomach wall	9.2E-11	3.7E-11	4.1E-11	4.7E-11	3.8E-09	8.4E-12	4.3E-11	2.1E-11	7.5E-11	1.5E-11	6.2E-11
SI + contents	1.1E-10	4.2E-11	4.7E-11	5.4E-11	2.6E-09	4.2E-12	3.2E-12	1.1E-12	3.1E-12	1.0E-12	1.4E-12
LLI wall	1.4E-10	1.0E-10	1.3E-10	1.6E-10	2.7E-09	2.3E-12	3.2E-12	1.1E-12	3.2E-12	1.1E-12	1.4E-12
LLI wall	5.5E-11	1.2E-10	1.6E-10	1.9E-10	2.7E-09	3.8E-13	3.2E-12	1.1E-12	2.9E-12	8.0E-13	1.3E-12
Liver	7.7E-12	3.3E-11	1.7E-11	1.4E-11	4.3E-09	2.6E-14	3.2E-12	1.1E-12	4.2E-12	2.3E-12	1.6E-12
Kidneys	7.1E-12	1.4E-11	7.7E-12	6.8E-12	3.1E-09	2.6E-14	3.2E-12	1.1E-12	3.5E-12	1.4E-12	1.5E-12
Bladder wall	6.2E-12	1.3E-11	6.6E-12	5.7E-12	2.3E-09	2.6E-14	3.2E-12	1.1E-12	2.7E-12	7.8E-13	1.3E-12
Muscle	6.5E-12	1.2E-11	7.2E-12	6.4E-12	3.1E-09	2.6E-14	3.2E-12	1.1E-12	3.3E-12	1.6E-12	1.4E-12
Bone Surface	5.8E-12	1.2E-11	6.2E-12	5.2E-12	3.4E-09	3.2E-14	3.2E-12	1.1E-12	3.0E-12	1.3E-12	1.5E-12
Skin	5.3E-12	9.8E-12	5.0E-12	4.1E-12	1.9E-09	2.6E-14	3.2E-12	1.1E-12	2.6E-12	1.1E-12	1.3E-12
Spleen	7.9E-12	1.4E-11	9.2E-12	8.4E-12	4.0E-09	2.6E-14	3.2E-12	1.1E-12	4.9E-12	2.3E-12	1.7E-12
Uterus	7.8E-12	1.4E-11	7.9E-12	7.4E-12	2.7E-09	2.6E-14	3.2E-12	1.1E-12	3.0E-12	9.0E-13	1.3E-12
Pancreas	9.2E-12	4.6E-11	2.3E-11	1.9E-11	4.2E-09	2.6E-14	3.2E-12	1.1E-12	6.1E-12	2.6E-12	2.0E-12
Total Body	1.2E-11	1.6E-11	1.3E-11	1.3E-11	3.3E-09	1.2E-12	5.3E-12	3.7E-12	5.7E-12	4.0E-12	3.7E-12
Remainder					4.3E-09						
Nuclide	Rb-89	Sr-89	Sr-89	Sr-90	Sr-90	Y-90	Y-90	Y-91m	Y-91m	Y-91	Y-91
f1	1.0	0.3	0.01	0.3	0.01	0.0001	0.0001	0.0001	0.0001	0.0001	0.0001
Class	0	0	Y	0	Y	W	Y	W	Y	W	Y
ALI (Bq)	4.9E+09	3.7E+07	5.2E+06	7.3E+05	1.4E+05	2.5E+07	2.3E+07	9.0E+09	6.0E+09	7.0E+06	4.2E+06
DAC (Bq/m ³)	2.1E+06	1.5E+04	2.2E+03	3.0E+02	5.8E+01	1.0E+04	9.5E+03	3.8E+06	2.5E+06	2.9E+03	1.8E+03
Lungs	6.9E-11	2.2E-09	8.1E-08	1.1E-09	3.0E-06	8.9E-09	9.3E-09	4.2E-11	7.0E-11	5.3E-08	9.9E-08
Thyroid	1.6E-12	3.9E-10	7.5E-12	0.0E-01	0.0E-01	9.4E-12	5.1E-13	6.0E-13	5.0E-13	7.3E-11	6.4E-12
Testes	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01
Ovaries	1.3E-12	3.9E-10	7.5E-12	0.0E-01	0.0E-01	9.4E-12	5.1E-13	4.1E-13	3.2E-13	7.3E-11	6.1E-12
Red Marrow	2.3E-12	2.8E-09	5.3E-11	2.3E-07	2.2E-08	2.7E-10	1.5E-11	2.7E-12	7.1E-13	3.5E-09	2.0E-10
Stomach wall	3.0E-11	3.9E-10	7.5E-12	0.0E-01	0.0E-01	3.8E-10	4.3E-10	2.4E-12	2.7E-12	3.9E-10	3.4E-10
SI + contents	1.5E-12	3.9E-10	7.5E-12	0.0E-01	0.0E-01	8.9E-10	1.0E-09	1.6E-12	1.7E-12	8.7E-10	8.4E-10
LLI wall	1.5E-12	1.5E-09	5.0E-09	6.6E-10	5.7E-09	4.7E-09	5.4E-09	3.8E-12	4.0E-12	4.9E-09	5.0E-09
LLI wall	1.3E-12	3.7E-09	1.4E-08	2.8E-09	2.0E-08	1.1E-08	1.3E-08	8.5E-12	8.8E-12	1.4E-08	1.5E-08
Liver	2.2E-12	3.9E-10	7.5E-12	0.0E-01	0.0E-01	2.7E-10	1.5E-11	3.2E-12	1.2E-12	3.5E-09	2.0E-10
Kidneys	1.7E-12	3.9E-10	7.4E-12	0.0E-01	0.0E-01	9.4E-12	5.1E-13	6.6E-13	5.7E-13	7.3E-11	6.3E-12
Bladder wall	1.2E-12	3.9E-10	7.4E-12	0.0E-01	0.0E-01	9.4E-12	5.1E-13	2.6E-13	1.2E-13	7.2E-11	4.8E-12
Muscle	1.7E-12	3.9E-10	7.5E-12	0.0E-01	0.0E-01	9.4E-12	5.1E-13	6.9E-13	6.1E-13	7.3E-11	6.8E-12
Bone Surface	2.4E-12	7.6E-09	1.4E-10	6.9E-07	6.6E-08	2.7E-10	1.5E-11	2.6E-12	5.5E-13	3.6E-09	2.0E-10
Skin	1.3E-12	3.9E-10	7.4E-12	0.0E-01	0.0E-01	9.4E-12	5.1E-13	3.8E-13	3.0E-13	7.2E-11	5.5E-12
Spleen	2.6E-12	3.9E-10	7.5E-12	0.0E-01	0.0E-01	9.4E-12	5.1E-13	1.1E-12	1.1E-12	7.4E-11	8.1E-12
Uterus	1.3E-12	3.9E-10	7.4E-12	0.0E-01	0.0E-01	9.4E-12	5.1E-13	3.6E-13	2.5E-13	7.2E-11	5.2E-12
Pancreas	3.2E-12	3.9E-10	7.5E-12	0.0E-01	0.0E-01	9.4E-12	5.1E-13	1.5E-12	1.5E-12	7.4E-11	8.6E-12
Total Body	2.8E-12	1.2E-09	1.2E-09	6.9E-08	4.9E-08	2.1E-10	1.9E-10	1.5E-12	1.6E-12	1.3E-09	1.5E-09
Remainder		1.2E-09		6.9E-08							

TABLE 3.10 Specific committed dose equivalents (Sv/Bq) for selected inhaled radionuclides and their daughters.
Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks.

Nuclide	Y-92	Y-92	Y-93	Y-93	Zr-95	Zr-95	Zr-95	Zr-97	Zr-97	Nb-95
f1	0.0001	0.0001	0.0001	0.0001	0.002	0.002	0.002	0.002	0.001	0.01
Class	W	Y	W	Y	W	Y	W	Y	W	W
ALI (Bq)	3.2E+08	2.9E+08	1.1E+08	8.9E+07	1.4E+07	1.0E+07	4.9E+06	5.2E+07	4.7E+07	7.4E+07
OAC (Bq/m ³)	1.3E+05	1.2E+05	4.4E+04	3.7E+04	6.0E+03	4.3E+03	2.0E+03	2.2E+04	2.0E+04	3.1E+04
Lungs	1.2E-09	1.2E-09	2.4E-09	2.5E-09	1.8E-08	4.0E-08	2.2E-09	3.9E-09	4.0E-09	2.1E-09
Thyroid	3.7E-12	1.1E-12	5.1E-12	9.2E-13	7.8E-10	1.2E-09	1.4E-09	3.7E-11	2.3E-11	9.6E-11
Testes	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01
Ovaries	4.9E-12	2.6E-12	8.7E-12	5.3E-12	8.4E-10	5.7E-10	1.9E-09	1.7E-10	1.8E-10	1.8E-10
Red Marrow	1.3E-11	2.1E-12	4.1E-11	4.0E-12	3.2E-09	1.3E-09	1.3E-08	1.4E-10	6.3E-11	4.8E-10
Stomach wall	1.4E-10	1.7E-10	2.4E-10	2.9E-10	1.1E-09	1.8E-09	1.1E-09	4.1E-10	4.7E-10	3.0E-10
SI + contents	1.9E-10	2.4E-10	4.7E-10	5.7E-10	9.8E-10	8.4E-10	1.7E-09	8.8E-10	1.0E-09	6.2E-10
ULI wall	3.2E-10	4.0E-10	1.5E-09	1.8E-09	1.9E-09	1.9E-09	1.8E-09	3.0E-09	3.5E-09	2.0E-09
LLI wall	1.7E-10	2.0E-10	1.6E-09	2.0E-09	4.1E-09	3.9E-09	3.0E-09	4.2E-09	5.1E-09	2.8E-09
Liver	1.3E-11	2.8E-12	4.1E-11	4.3E-12	1.2E-09	2.1E-09	1.3E-09	7.4E-11	6.4E-11	1.2E-10
Kidneys	4.2E-12	1.7E-12	5.9E-12	1.9E-12	8.1E-10	9.6E-10	1.9E-09	6.1E-11	5.1E-11	1.2E-10
Bladder wall	3.5E-12	8.9E-13	5.7E-12	1.8E-12	3.8E-10	2.1E-10	1.1E-09	6.1E-11	5.3E-11	1.2E-10
Muscle	4.1E-12	1.5E-12	5.8E-12	1.7E-12	9.3E-10	1.2E-09	1.9E-09	5.8E-11	4.7E-11	1.1E-10
Bone Surface	1.2E-11	1.5E-12	4.0E-11	3.1E-12	2.2E-08	2.3E-09	1.0E-07	1.2E-10	3.5E-11	5.1E-10
Skin	3.4E-12	8.1E-13	5.0E-12	9.4E-13	5.8E-10	6.3E-10	1.5E-09	3.3E-11	2.1E-11	8.6E-11
Spleen	4.9E-12	2.4E-12	6.4E-12	2.4E-12	1.1E-09	1.9E-09	1.6E-09	7.2E-11	6.1E-11	1.2E-10
Uterus	4.3E-12	1.8E-12	6.7E-12	2.9E-12	5.1E-10	2.8E-10	1.5E-09	8.9E-11	8.6E-11	1.4E-10
Pancreas	5.4E-12	3.0E-12	6.9E-12	2.9E-12	1.3E-09	2.2E-09	1.7E-09	8.6E-11	7.7E-11	1.3E-10
Total Body	2.5E-11	2.3E-11	5.7E-11	5.4E-11	1.4E-09	1.8E-09	3.1E-09	1.5E-10	1.4E-10	1.9E-10
Remainder										4.8E-10

Nuclide	Nb-95	Mo-99	Mo-99	Tc-99m	Tc-99m	Tc-101	Tc-101	Ru-103	Ru-103	Ru-103	Ru-105
f1	0.01	0.05	0.8	0.8	0.8	0.8	0.8	0.05	0.05	0.05	0.05
Class	Y	Y	W	W	W	W	W	W	W	Y	W
ALI (Bq)	4.1E+07	5.0E+07	9.4E+07	9.6E+09	6.4E+09	1.4E+10	1.2E+10	3.8E+07	6.3E+07	2.5E+07	5.7E+08
OAC (Bq/m ³)	1.7E+04	2.1E+04	3.9E+04	4.0E+06	2.7E+06	5.8E+06	5.1E+06	1.6E+04	2.6E+04	1.0E+04	2.4E+05
Lungs	8.3E-09	4.3E-09	1.2E-09	3.0E-11	2.2E-11	3.0E-11	2.8E-11	9.5E-09	1.0E-09	1.5E-08	5.2E-10
Thyroid	3.6E-10	1.5E-11	1.2E-10	2.1E-11	5.0E-11	7.4E-14	7.1E-14	2.8E-10	6.0E-10	2.6E-08	6.2E-12
Testes	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01
Ovaries	4.3E-10	9.7E-11	1.3E-10	1.7E-12	2.8E-12	1.4E-14	4.6E-14	4.0E-10	7.4E-10	3.2E-10	1.5E-11
Red Marrow	4.4E-10	5.1E-11	3.4E-10	2.4E-12	3.4E-12	1.0E-13	1.2E-13	3.5E-10	6.7E-10	3.3E-10	9.1E-12
Stomach wall	6.3E-10	2.3E-10	2.2E-10	9.5E-12	1.6E-11	1.7E-12	1.2E-11	4.9E-10	7.2E-10	5.1E-10	6.8E-11
SI + contents	5.2E-10	5.2E-10	1.8E-10	3.5E-12	4.9E-12	2.0E-13	1.6E-12	5.5E-10	8.5E-10	4.7E-10	1.0E-10
ULI wall	9.9E-10	2.4E-09	3.4E-10	5.5E-12	6.9E-12	3.7E-14	1.4E-13	1.3E-09	1.1E-09	1.3E-09	2.2E-10
LLI wall	1.9E-09	5.5E-09	5.8E-10	3.7E-12	5.0E-12	1.2E-14	4.6E-14	3.0E-09	1.7E-09	3.0E-09	2.1E-10
Liver	6.7E-10	1.1E-10	1.9E-09	2.8E-12	4.0E-12	1.8E-13	2.0E-13	4.6E-10	6.8E-10	5.1E-10	1.2E-11
Kidneys	3.5E-10	1.0E-10	1.9E-09	1.6E-12	2.6E-12	7.7E-14	1.2E-13	3.1E-10	6.9E-10	2.7E-10	8.8E-12
Bladder wall	1.4E-10	3.3E-11	1.3E-10	9.0E-13	1.9E-12	6.2E-15	1.8E-14	2.4E-10	7.1E-10	1.3E-10	6.4E-12
Muscle	4.1E-10	2.8E-11	1.3E-10	1.5E-12	2.2E-12	9.9E-14	1.1E-13	3.2E-10	6.1E-10	3.2E-10	8.1E-12
Bone Surface	5.1E-10	4.1E-11	5.3E-10	1.8E-12	2.8E-12	8.3E-14	1.1E-13	2.8E-10	6.2E-10	2.4E-10	6.5E-12
Skin	1.1E-10	1.4E-11	1.1E-10	7.4E-13	1.2E-12	4.7E-14	5.2E-14	2.0E-10	4.6E-10	1.7E-10	4.9E-12
Spleen	6.3E-10	3.6E-11	1.4E-10	2.8E-12	3.8E-12	1.8E-13	2.9E-13	4.3E-10	7.0E-10	4.7E-10	1.2E-11
Uterus	1.9E-10	4.5E-11	1.3E-10	1.3E-12	2.5E-12	1.2E-14	4.6E-14	2.6E-10	7.0E-10	1.6E-10	9.9E-12
Pancreas	7.1E-10	4.3E-11	1.7E-10	3.6E-12	5.1E-12	2.2E-13	4.3E-13	5.1E-10	6.9E-10	6.0E-10	1.4E-11
Total Body	5.1E-10	1.2E-10	2.2E-10	2.1E-12	2.7E-12	5.3E-13	5.5E-13	4.6E-10	6.2E-10	5.3E-10	1.7E-11
Remainder					5.1E-12				7.1E-10		

TABLE 3.11 Specific committed dose equivalents (Sv/Bq) for selected inhaled radionuclides and their daughters.
Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks.

Nuclide	Ru-105	Ru-105	Ru-106	Ru-106	Ru-106	Ag-110m	Ag-110m	Ag-110m	Te-125m	Te-125m	Te-127m
f1	0.05	0.05	0.05	0.05	0.05	0.05	0.05	0.05	0.2	0.2	0.2
Class	Y	0	W	Y	0	W	Y	0	0	0	0
ALI (Bq)	4.8E+08	5.6E+08	2.0E+06	4.0E+05	3.3E+06	7.5E+06	3.5E+06	5.2E+06	1.4E+07	2.4E+07	9.6E+06
DAC (Bq/m ³)	2.0E+05	2.4E+05	8.3E+02	1.7E+02	1.4E+03	3.1E+03	1.5E+03	2.2E+03	5.7E+03	9.9E+03	4.0E+03
Lungs	5.5E-10	3.6E-10	2.1E-07	1.0E-06	1.8E-08	3.1E-08	1.2E-07	8.0E-09	5.2E-10	1.2E-08	8.9E-10
Thyroid	4.0E-12	1.4E-11	4.0E-09	1.7E-09	1.4E-08	2.0E-09	6.3E-09	1.7E-09	1.1E-10	4.2E-11	2.4E-10
Testes	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01
Ovaries	1.5E-11	2.6E-11	4.0E-09	1.3E-09	1.4E-08	2.3E-09	2.4E-09	3.2E-09	1.3E-10	8.2E-11	2.5E-10
Red Marrow	7.4E-12	1.8E-11	4.1E-09	1.8E-09	1.4E-08	2.9E-09	6.7E-09	4.0E-09	3.4E-09	1.3E-09	1.4E-08
Stomach wall	8.0E-11	8.8E-11	5.0E-09	2.9E-09	1.4E-08	4.5E-09	1.0E-08	6.5E-09	1.4E-10	1.4E-10	2.6E-10
SI + contents	1.2E-10	1.3E-10	6.0E-09	3.4E-09	1.5E-08	3.4E-09	3.4E-09	6.0E-09	1.7E-10	2.1E-10	2.9E-10
ULI wall	2.6E-10	2.4E-10	1.6E-08	1.4E-08	1.8E-08	5.1E-09	5.2E-09	8.4E-09	4.0E-10	9.1E-10	7.2E-10
LLI wall	2.5E-10	1.9E-10	3.9E-08	3.7E-08	2.5E-08	5.8E-09	5.8E-09	3.9E-09	9.4E-10	2.5E-09	1.9E-09
Liver	9.9E-12	2.0E-11	4.2E-09	2.3E-09	1.4E-08	2.5E-08	1.8E-08	8.0E-08	1.1E-10	8.0E-11	2.4E-10
Kidneys	7.0E-12	1.8E-11	4.1E-09	1.7E-09	1.4E-08	4.5E-09	6.2E-09	1.1E-08	1.1E-10	4.5E-11	2.4E-10
Bladder wall	4.4E-12	1.7E-11	4.0E-09	1.2E-09	1.4E-08	1.1E-09	1.0E-09	2.2E-09	1.1E-10	4.4E-11	2.4E-10
Muscle	6.4E-12	1.6E-11	4.0E-09	1.8E-09	1.4E-08	2.9E-09	7.1E-09	4.1E-09	1.2E-10	7.4E-11	2.4E-10
Bone Surface	4.4E-12	1.5E-11	4.0E-09	1.6E-09	1.4E-08	2.1E-09	5.2E-09	3.0E-09	3.7E-08	1.3E-08	5.2E-08
Skin	3.0E-12	1.2E-11	3.9E-09	1.4E-09	1.3E-08	1.6E-09	3.7E-09	2.4E-09	1.1E-10	4.4E-11	2.4E-10
Spleen	1.0E-11	2.0E-11	4.2E-09	2.2E-09	1.4E-08	3.8E-09	1.1E-08	4.2E-09	1.2E-10	7.6E-11	2.4E-10
Uterus	8.6E-12	2.1E-11	4.0E-09	1.2E-09	1.4E-08	1.5E-09	1.5E-09	2.9E-09	1.1E-10	4.7E-11	2.4E-10
Pancreas	1.3E-11	2.3E-11	4.2E-09	2.5E-09	1.4E-08	6.7E-09	1.3E-08	1.3E-08	1.1E-10	6.6E-11	2.4E-10
Total Body	1.7E-11	2.4E-11	7.1E-09	1.7E-08	1.4E-08	3.7E-09	8.4E-09	5.9E-09	4.3E-10	3.6E-10	1.4E-09
Remainder					1.4E-08	6.7E-09		1.3E-08			
Nuclide	Ta-127m	Te-127	Te-127	Te-129m	Te-129m	Te-129	Te-129	Te-131m	Te-131	Te-132	1-125
f1	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	1.0
Class	W	0	W	0	W	0	W	0	0	0	0
ALI (Bq)	9.5E+06	8.4E+08	6.4E+08	2.4E+07	9.1E+06	2.4E+09	2.7E+09	1.5E+07	1.8E+08	8.4E+06	2.4E+06
DAC (Bq/m ³)	4.0E+03	3.5E+05	2.7E+05	9.8E+03	3.8E+03	9.8E+05	1.1E+06	6.3E+03	7.7E+04	3.5E+03	1.0E+03
Lungs	3.3E-08	2.8E-10	4.3E-10	2.2E-09	4.0E-08	1.3E-10	1.5E-10	5.9E-10	2.5E-10	6.2E-10	1.1E-10
Thyroid	9.6E-11	6.5E-12	1.8E-12	3.9E-10	1.5E-10	1.6E-12	5.1E-13	3.3E-08	2.7E-09	6.0E-08	2.1E-07
Testes	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01
Ovaries	1.1E-10	6.6E-12	2.0E-12	4.1E-10	1.8E-10	1.8E-12	5.0E-13	1.6E-10	5.5E-12	3.4E-10	1.3E-11
Red Marrow	5.3E-09	1.4E-11	4.0E-12	8.6E-09	3.0E-09	2.0E-12	6.1E-13	9.4E-11	5.5E-12	3.5E-10	3.6E-11
Stomach wall	2.2E-10	4.2E-11	4.5E-11	4.7E-10	4.5E-10	5.2E-11	1.6E-11	9.9E-11	8.7E-11	3.4E-10	1.9E-11
SI + contents	3.5E-10	6.4E-11	7.1E-11	6.0E-10	7.8E-10	3.6E-11	1.1E-11	2.5E-10	7.6E-11	3.8E-10	1.4E-11
ULI wall	1.8E-09	1.8E-10	2.1E-10	1.7E-09	3.9E-09	2.5E-11	7.5E-12	5.1E-10	7.5E-11	3.6E-10	1.4E-11
LLI wall	5.6E-09	1.9E-10	2.2E-10	4.2E-09	1.1E-08	6.1E-12	1.8E-12	7.9E-10	5.1E-12	8.7E-10	1.3E-11
Liver	1.1E-10	6.5E-12	1.9E-12	3.9E-10	1.8E-10	1.8E-12	6.0E-13	9.5E-11	5.6E-12	3.5E-10	1.5E-11
Kidneys	9.7E-11	6.5E-12	1.9E-12	4.0E-10	1.6E-10	1.7E-12	5.2E-13	8.6E-11	5.2E-12	3.5E-10	1.2E-11
Bladder wall	9.3E-11	6.5E-12	1.9E-12	3.9E-10	1.5E-10	1.6E-12	4.7E-13	9.4E-11	4.5E-12	3.7E-10	1.3E-11
Muscle	1.1E-10	6.5E-12	1.9E-12	4.0E-10	1.7E-10	1.7E-12	5.4E-13	8.3E-11	5.2E-12	3.3E-10	8.9E-11
Bone Surface	2.0E-08	1.4E-11	4.1E-12	1.9E-08	6.8E-09	2.0E-12	6.2E-13	7.4E-11	4.9E-12	3.3E-10	3.6E-11
Skin	9.5E-11	6.4E-12	1.8E-12	3.9E-10	1.5E-10	1.6E-12	4.8E-13	5.4E-11	4.2E-12	2.5E-10	4.1E-11
Spleen	1.1E-10	6.5E-12	1.9E-12	4.0E-10	1.8E-10	1.8E-12	6.1E-13	9.1E-11	6.0E-12	3.5E-10	1.5E-11
Uterus	9.5E-11	6.5E-12	1.9E-12	4.0E-10	1.5E-10	1.7E-12	4.9E-13	1.2E-10	5.4E-12	3.9E-10	1.3E-11
Pancreas	1.1E-10	6.6E-12	2.0E-12	4.0E-10	1.8E-10	1.9E-12	6.5E-13	9.5E-11	6.9E-12	3.5E-10	1.5E-11
Total Body	1.1E-09	1.3E-11	1.0E-11	1.1E-09	1.0E-09	6.1E-12	2.9E-12	1.0E-10	1.0E-11	3.5E-10	1.3E-10
Remainder											

TABLE 3.12 Specific committed dose equivalents (Sv/Bq) for selected inhaled radionuclides and their daughters.
Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks.

Nuclide	I-130	I-131	I-133	I-134	I-135	Ca-134	Ca-136	Ca-137	Ca-138	Ba-139	Ba-140
f1	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	0.1	0.1
Class	0	0	0	0	0	0	0	0	0	0	0
ALI (Bq)	2.6E+07	1.6E+06	1.1E+07	1.7E+09	6.4E+07	4.1E+06	2.6E+07	6.1E+06	2.1E+09	1.1E+09	5.2E+07
OAC (Bq/m ³)	1.2E+04	7.5E+02	4.6E+03	7.3E+05	2.7E+04	1.7E+03	1.1E+04	2.6E+03	8.7E+05	4.8E+05	2.2E+04
Lungs	5.9E-10	6.5E-10	8.2E-10	1.4E-10	4.4E-10	1.2E-08	2.3E-09	8.5E-09	1.6E-10	2.5E-10	1.6E-09
Thyroid	1.8E-08	2.8E-07	4.5E-08	2.6E-10	7.8E-09	1.1E-08	1.7E-09	7.6E-09	3.6E-12	2.4E-12	2.4E-10
Testes	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01
Ovaries	2.7E-11	2.2E-11	1.9E-11	4.2E-12	1.7E-11	1.1E-08	1.7E-09	7.8E-09	3.3E-12	2.5E-12	4.2E-10
Red Marrow	4.5E-11	5.7E-11	2.7E-11	6.0E-12	2.2E-11	1.2E-08	1.9E-09	8.0E-09	4.0E-12	2.5E-12	1.4E-09
Stomach wall	1.2E-10	7.1E-11	1.0E-10	7.0E-11	1.0E-10	1.2E-08	2.0E-09	8.3E-09	7.8E-11	9.3E-11	2.6E-10
SI + contents	3.4E-11	2.3E-11	2.2E-11	5.5E-12	1.9E-11	1.4E-08	2.1E-09	8.6E-09	3.7E-12	7.3E-11	5.1E-10
ULI wall	3.5E-11	2.3E-11	2.2E-11	5.7E-12	1.9E-11	1.3E-08	2.0E-09	8.7E-09	3.9E-12	5.9E-11	1.4E-09
LLI wall	3.1E-11	2.2E-11	2.1E-11	4.6E-12	1.6E-11	1.4E-08	2.1E-09	8.8E-09	3.4E-12	1.5E-11	4.3E-09
Liver	5.4E-11	3.5E-11	2.9E-11	8.4E-12	2.6E-11	1.3E-08	2.0E-09	8.3E-09	5.2E-12	2.5E-12	2.7E-10
Kidneys	4.0E-11	2.6E-11	2.4E-11	6.6E-12	2.2E-11	1.3E-08	2.0E-09	8.3E-09	4.3E-12	2.4E-12	2.8E-10
Bladder wall	3.0E-11	2.1E-11	2.0E-11	4.5E-12	1.7E-11	1.3E-08	2.1E-09	8.6E-09	3.1E-12	2.4E-12	3.0E-10
Muscle	4.8E-11	7.2E-11	2.9E-11	6.1E-12	2.3E-11	1.1E-08	1.7E-09	7.5E-09	4.0E-12	2.4E-12	2.8E-10
Bone Surface	4.0E-11	5.2E-11	2.5E-11	5.3E-12	2.0E-11	1.1E-08	1.7E-09	7.6E-09	3.6E-12	2.4E-12	2.6E-09
Skin	3.1E-11	4.8E-11	2.2E-11	4.1E-12	1.7E-11	7.6E-09	1.2E-09	6.3E-09	3.1E-12	2.4E-12	2.2E-10
Spleen	5.6E-11	3.5E-11	2.9E-11	1.0E-11	2.6E-11	1.3E-08	2.0E-09	8.3E-09	6.2E-12	2.5E-12	2.6E-10
Uterus	3.1E-11	2.2E-11	2.0E-11	5.0E-12	1.9E-11	1.4E-08	2.4E-09	8.7E-09	3.4E-12	2.5E-12	3.0E-10
Pancreas	6.4E-11	3.6E-11	3.2E-11	1.3E-11	3.4E-11	1.2E-08	1.9E-09	8.1E-09	8.0E-12	2.6E-12	3.0E-10
Total Body	5.9E-11	1.5E-10	5.2E-11	8.2E-12	3.1E-11	1.1E-08	1.7E-09	7.5E-09	6.4E-12	7.0E-12	4.1E-10
Remainder						1.4E-08	2.4E-09	8.7E-09			
Nuclide	Ba-141	La-140	La-140	La-142	La-142	Ca-141	Ca-141	Ca-143	Ca-143	Ca-144	Ca-144
f1	0.1	0.001	0.001	0.001	0.001	0.0003	0.0003	0.0003	0.0003	0.0003	0.0003
Class	0	0	0	0	0	0	0	0	0	0	0
ALI (Bq)	2.6E+09	5.5E+07	4.3E+07	8.0E+08	1.2E+09	2.6E+07	2.2E+07	6.6E+07	5.9E+07	9.5E+05	5.3E+05
OAC (Bq/m ³)	1.1E+06	2.3E+04	1.6E+04	3.4E+05	4.6E+05	1.1E+04	9.3E+03	2.7E+04	2.4E+04	4.0E+02	2.2E+02
Lungs	1.1E-10	1.7E-09	4.2E-09	3.1E-10	3.6E-10	1.1E-08	1.7E-08	3.6E-09	3.9E-09	1.6E-07	7.6E-07
Thyroid	1.3E-12	1.2E-10	6.9E-11	9.0E-12	5.0E-12	4.6E-11	2.6E-11	1.2E-11	6.0E-12	1.9E-09	2.9E-10
Testes	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01
Ovaries	1.4E-12	3.6E-10	4.5E-10	1.7E-11	6.2E-12	8.5E-11	5.6E-11	6.9E-11	7.4E-11	1.9E-09	2.3E-10
Red Marrow	1.5E-12	4.4E-10	2.1E-10	1.4E-11	7.0E-12	4.1E-10	9.0E-11	7.6E-11	2.9E-11	2.5E-08	2.7E-09
Stomach wall	3.9E-11	3.9E-10	4.7E-10	1.2E-10	5.1E-11	1.9E-10	1.6E-10	1.9E-10	2.1E-10	2.7E-09	1.1E-09
SI + contents	2.3E-11	6.5E-10	9.7E-10	1.1E-10	4.3E-11	3.1E-10	2.9E-10	4.3E-10	4.9E-10	3.8E-09	2.1E-09
ULI wall	3.1E-11	1.6E-09	2.9E-09	1.1E-10	4.1E-11	1.4E-09	1.5E-09	1.8E-09	2.1E-09	1.3E-08	1.2E-08
LLI wall	1.7E-11	2.6E-09	5.4E-09	3.4E-11	1.2E-11	3.6E-09	4.1E-09	3.7E-09	4.3E-09	3.4E-08	3.4E-08
Liver	1.7E-12	3.5E-09	7.6E-10	3.9E-11	1.6E-11	3.4E-09	2.6E-10	5.1E-10	4.8E-11	2.5E-07	2.5E-08
Kidneys	1.5E-12	3.4E-10	1.7E-10	1.4E-11	6.3E-12	9.6E-11	3.2E-11	2.4E-11	1.5E-11	2.1E-09	3.1E-10
Bladder wall	1.2E-12	1.9E-10	1.7E-10	9.7E-12	3.2E-12	4.7E-11	1.8E-11	2.3E-11	2.0E-11	1.9E-09	2.0E-10
Muscle	1.4E-12	2.0E-10	1.5E-10	1.2E-11	6.4E-12	7.1E-11	4.5E-11	2.2E-11	1.6E-11	1.9E-09	3.4E-10
Bone Surface	1.3E-12	4.0E-10	1.4E-10	1.1E-11	5.5E-12	3.8E-09	2.5E-10	7.8E-11	1.6E-11	4.5E-08	4.7E-09
Skin	1.2E-12	1.4E-10	7.6E-11	7.8E-12	3.9E-12	4.4E-11	1.8E-11	1.2E-11	6.4E-12	1.9E-09	2.5E-10
Spleen	2.0E-12	2.1E-10	1.7E-10	1.7E-11	1.0E-11	2.9E-09	2.2E-10	4.4E-10	4.3E-11	2.2E-07	2.2E-08
Uterus	1.3E-12	2.3E-10	2.3E-10	1.5E-11	5.3E-12	5.6E-11	2.6E-11	3.2E-11	3.0E-11	1.9E-09	2.1E-10
Pancreas	2.4E-12	3.9E-10	2.4E-10	2.3E-11	1.2E-11	1.4E-10	8.2E-11	3.5E-11	2.5E-11	2.2E-09	4.7E-10
Total Body	3.4E-12	3.4E-10	2.5E-10	1.6E-11	1.2E-11	3.7E-10	3.1E-10	1.1E-10	9.4E-11	1.4E-08	1.3E-08
Remainder											

TABLE 3.13 Specific committed dose equivalents (Sv/Bq) for selected inhaled radionuclides and their daughters.
Annual limits of intake are the lesser of the values for stochastic and non-stochastic risks.

=====								
Nuclide	Pr-143	Pr-145	Pr-144	Pr-144	Nd-147	Nd-147	W-187	Np-239
f1	0.0003	0.0005	0.0005	0.0003	0.0003	0.0005	0.5	0.01
Class	W	Y	W	Y	W	Y	0	W
ALI (Bq)	5.0E+07	2.5E+07	4.7E+09	4.4E+09	5.5E+07	2.9E+07	5.5E+08	9.2E+07
DAC (Bq/m ³)	1.2E+04	1.0E+04	2.0E+06	1.8E+06	1.5E+04	1.2E+04	1.5E+05	5.8E+04

Lungs	1.1E-08	1.5E-08	8.9E-11	9.4E-11	8.4E-09	1.1E-08	6.0E-10	2.2E-09
Thyroid	1.6E-18	1.7E-18	8.0E-15	8.5E-15	1.9E-11	1.8E-11	4.5E-12	5.8E-12
Teates	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01	0.0E-01
Ovaries	4.3E-18	4.4E-18	2.2E-15	2.4E-15	8.0E-11	8.5E-11	3.0E-11	7.5E-11
Red Marrow	2.7E-10	1.5E-11	1.6E-15	1.8E-14	5.0E-10	9.2E-11	2.7E-11	1.4E-10
Stomach wall	1.5E-10	1.7E-10	4.2E-12	5.4E-12	1.8E-10	2.0E-10	7.0E-11	1.2E-10
SI + contents	3.7E-10	4.1E-10	9.8E-15	1.3E-12	4.0E-10	4.5E-10	1.2E-10	2.8E-10
ULI wall	2.2E-09	2.4E-09	1.8E-15	2.4E-15	1.9E-09	2.2E-09	4.0E-10	1.2E-09
LLI wall	6.1E-09	6.8E-09	1.0E-14	1.3E-14	5.2E-09	5.8E-09	6.6E-10	2.7E-09
Liver	2.2E-09	1.2E-10	5.4E-15	3.5E-14	1.6E-09	1.8E-10	5.2E-11	5.8E-10
Kidneys	4.9E-10	2.7E-11	2.6E-13	2.2E-14	4.0E-11	2.6E-11	8.5E-11	1.6E-11
Bladder wall	1.4E-18	1.5E-18	1.1E-15	1.1E-15	2.4E-11	2.4E-11	8.7E-12	1.7E-11
Muscle	2.5E-18	2.2E-18	9.9E-15	1.0E-14	5.8E-11	3.5E-11	8.8E-12	1.5E-11
Bone Surface	2.7E-10	1.5E-11	1.5E-13	1.6E-14	2.5E-09	5.2E-10	9.7E-11	1.5E-09
Skin	1.5E-18	1.1E-18	5.4E-15	5.7E-15	1.7E-11	1.4E-11	4.0E-12	5.6E-12
Spleen	5.3E-18	5.2E-18	1.7E-14	1.8E-14	4.7E-11	4.8E-11	7.5E-11	1.8E-11
Uterus	1.9E-18	1.9E-18	2.1E-15	2.5E-15	5.5E-11	3.5E-11	1.5E-11	2.4E-11
Pancreas	5.0E-18	3.8E-18	1.9E-14	2.1E-14	7.0E-11	5.8E-11	1.7E-11	2.5E-11
Total Body	2.6E-10	2.2E-10	1.5E-12	1.4E-12	2.6E-10	2.2E-10	2.5E-11	7.9E-11
Remainder								
=====								

TABLE 3.14 Comparison of specific committed dose equivalents (Sv/Bq) to the results in ICRP Publication 30 for selected ingested radionuclides and their daughters. Values reported are greater than or equal to 10 % of the maximum.

=====										
Nuclide	P-32		Mn-54		Co-60		Ni-63		Zn-65	
f1	0.8		0.1		0.05		0.05		0.5	

Program	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE

Lungs			2.3E-10	2.3E-10	8.7E-10	8.8E-10	8.5E-11	8.5E-11	3.1E-09	3.1E-09
Thyroid									3.2E-09	3.2E-09
Testes										
Ovaries	6.5E-10	6.6E-10	9.5E-10	9.5E-10	3.2E-09	3.2E-09	8.5E-11	8.5E-11	3.5E-09	3.5E-09
Red Marrow	8.1E-09	7.8E-09	4.9E-10	5.1E-10	1.3E-09	1.3E-09	8.5E-11	8.5E-11	4.5E-09	4.9E-09
Stomach wall							1.0E-10	1.0E-10		
SI + contents			9.8E-10	9.9E-10	3.6E-09	3.6E-09	1.3E-10	1.3E-10	4.3E-09	4.3E-09
ULI well	3.0E-09	3.0E-09	1.4E-09	1.4E-09	5.7E-09	5.8E-09	3.6E-10	3.7E-10	4.2E-09	4.3E-09
LLI well	7.2E-09	7.2E-09	2.2E-09	2.2E-09	1.1E-08	1.1E-08	9.2E-10	9.2E-10	5.0E-09	5.0E-09
Liver			1.0E-09	1.0E-09	2.3E-09	2.3E-09				
Kidneys										
Bladder wall										
Muscle	6.5E-10	6.6E-10	2.8E-10	2.8E-10	1.1E-09	1.1E-09	8.5E-11	8.5E-11	3.3E-09	3.3E-09
Bone Surface	7.9E-09	7.8E-09							4.5E-09	4.5E-09
Skin										
Spleen										
Uterus										
Pancreas										
Total Body										
Remainder			5.0E-10	5.0E-10	2.1E-09	2.1E-09			4.8E-09	4.7E-09

Nuclide	Sr-89		Y-91m		Mo-99		Tc-99m		Ru-105	
f1	0.3		0.0001		0.8		0.8		0.05	

Program	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE

Lungs					1.9E-10	2.0E-10				
Thyroid							8.5E-11	8.4E-11		
Testes										
Ovaries			6.9E-12	6.9E-12	2.2E-10	2.3E-10	9.7E-12	9.5E-12	9.7E-11	9.5E-11
Red Marrow	3.2E-09	1.6E-09			5.3E-10	4.8E-10	6.3E-12	6.0E-12		
Stomach wall			4.9E-11	4.9E-11	6.7E-10	6.7E-10	7.2E-11	3.9E-11	5.0E-10	5.0E-10
SI + contents			3.1E-11	3.1E-11			2.2E-11	2.2E-11	7.9E-10	7.9E-10
ULI well	7.3E-09	7.2E-09	3.1E-11	3.1E-11	1.4E-09	1.5E-09	3.7E-11	3.6E-11	1.6E-09	1.6E-09
LLI well	2.1E-08	2.1E-08	2.4E-11	2.4E-11	3.1E-09	3.2E-09	2.5E-11	2.4E-11	1.3E-09	1.2E-09
Liver					2.7E-09	2.7E-09				
Kidneys					2.5E-09	2.7E-09				
Bladder wall										
Muscle					1.8E-10	1.9E-10	3.6E-12	3.3E-12		
Bone Surface	4.8E-09	4.3E-09			7.7E-10	7.6E-10				
Skin										
Spleen										
Uterus										
Pancreas										
Total Body										
Remainder			1.1E-11	1.1E-11			1.1E-11	9.3E-12		
=====										

TABLE 3.15 Comparison of specific committed dose equivalents (Sv/Bq) to the results in ICRP Publication 30 for selected ingested radionuclides and their daughters. Values reported are greater than or equal to 10 % of the maximum.

=====										
Nuclide	1e-127		1e-131m		I-131		I-135		Cs-137	
f1	0.2		0.2		1.0		1.0		1.0	

Program	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE

Lungs									1.3E-08	1.2E-08
Thyroid			4.3E-08	4.3E-08	4.8E-07	4.5E-07	1.8E-08	1.7E-08	1.3E-08	1.2E-08
Testes									1.3E-08	1.2E-08
Ovaries			7.3E-10	1.5E-11					1.4E-08	1.2E-08
Red Marrow									1.3E-08	1.3E-08
Stomach wall	2.4E-10	2.4E-10								
SI + contents	3.9E-10	3.9E-10								
ULI wall	1.2E-09	1.2E-09	4.6E-09	5.4E-10					1.4E-08	1.4E-08
LLI wall	1.3E-09	1.2E-09	8.2E-09	9.3E-12					1.4E-08	1.4E-08
Liver										
Kidneys										
Bladder wall										
Muscle									1.2E-08	1.2E-08
Bone Surface									1.3E-08	1.2E-08
Skin										
Spleen										
Uterus										
Pancreas										
Total Body										
Remainder									1.5E-08	1.4E-08

Nuclide	Be-141		Ce-143		Pr-144		W-187		Np-239	
f1	0.1		0.0003		0.0003		0.01		0.01	

Program	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE

Lungs										
Thyroid										
Testes										
Ovaries							2.6E-10	2.6E-10		
Red Marrow										
Stomach wall	3.9E-10	3.9E-10			4.1E-10	4.1E-10				
SI + contents	1.9E-10	1.8E-10	1.4E-09	1.4E-09	9.6E-11	9.6E-11	1.0E-09	1.0E-09	8.7E-10	8.2E-10
ULI wall	2.2E-10	2.2E-10	5.7E-09	5.9E-09			3.6E-09	3.6E-09	3.8E-09	3.7E-09
LLI wall	1.2E-10	1.1E-10	1.2E-08	1.2E-08			6.0E-09	5.9E-09	8.6E-09	8.1E-09
Liver										
Kidneys										
Bladder wall										
Muscle										
Bone Surface										
Skin										
Spleen										
Uterus										
Pancreas										
Total Body										
Remainder										
=====										

TABLE 3.16 Comparison of specific committed dose equivalents (Sv/Bq) to the results in ICRP Publication 30 for selected inhaled radionuclides and their daughters. Values reported are greater than or equal to 10 % of the maximum.

Nuclide	P-32		Cr-51		Mn-54		Fe-59		Co-60	
fl	0.8		0.01		0.1		0.1		0.05	
Class	D		D		W		D		W	
Program	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE
Lungs	2.5E-09	2.5E-09	3.8E-11	3.7E-11	6.7E-09	6.6E-09	3.5E-09	3.5E-09	3.6E-08	3.6E-08
Thyroid							2.9E-09	3.0E-09		
Testes										
Ovaries	4.8E-10	4.8E-10	2.7E-11	2.7E-11	7.1E-10	7.1E-10	3.3E-09	3.3E-09	4.0E-09	4.0E-09
Red Marrow	6.0E-09	5.8E-09	2.7E-11	3.1E-11	1.1E-09	1.1E-09	3.2E-09	3.2E-09	4.2E-09	4.2E-09
Stomach wall										
SI + contents			3.0E-11	3.0E-11						
ULI wall			3.8E-11	4.1E-11			4.1E-09	4.1E-09		
LLI wall	1.5E-09	1.5E-09	5.9E-11	6.8E-11			4.8E-09	4.8E-09	8.2E-09	8.1E-09
Liver					2.5E-09	2.5E-09	7.1E-09	7.1E-09	9.2E-09	9.2E-09
Kidneys										
Bladder wall										
Muscle	4.8E-10	4.8E-10	1.9E-11	1.9E-11	8.6E-10	8.6E-10	3.0E-09	3.0E-09	4.2E-09	4.2E-09
Bone Surface	5.8E-09	5.8E-09	2.7E-11	2.6E-11			2.9E-09	2.9E-09		
Skin										
Spleen							8.3E-09	8.5E-09		
Uterus										
Pancreas										
Total Body										
Remainder			2.5E-11	2.4E-11	1.8E-09	1.4E-09	4.7E-09	4.0E-09		

Nuclide	Sr-90		Y-93		Nb-95		Tc-99m		Ag-110m	
fl	0.3		0.0001		0.01		0.8		0.05	
Class	D		Y		W		D		W	
Program	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE
Lungs			2.5E-09	2.5E-09	5.5E-09	5.5E-09	2.3E-11	2.2E-11	3.2E-08	3.1E-08
Thyroid							5.0E-11	5.0E-11		
Testes										
Ovaries					4.8E-10	4.8E-10	2.8E-12	2.8E-12	2.3E-09	2.3E-09
Red Marrow	3.3E-07	2.3E-07			6.7E-10	6.7E-10	3.4E-12	3.4E-12		
Stomach wall							2.9E-11	1.6E-11		
SI + contents			5.7E-10	5.7E-10			4.8E-12	4.9E-12		
ULI wall			1.8E-09	1.8E-09			7.0E-12	6.9E-12		
LLI wall			2.0E-09	2.0E-09	1.9E-09	1.9E-09	5.0E-12	5.0E-12		
Liver									2.6E-08	2.5E-08
Kidneys										
Bladder wall										
Muscle							2.1E-12	2.2E-12	2.9E-09	2.9E-09
Bone Surface	7.3E-07	6.9E-07			2.4E-09	2.4E-09				
Skin										
Spleen										
Uterus										
Pancreas										
Total Body										
Remainder							5.0E-12	5.1E-12	6.9E-09	6.7E-09

TABLE 3.17 Comparison of specific committed dose equivalents (Sv/Bq) to the results in ICRP Publication 30 for selected inhaled radionuclides and their daughters. Values reported are greater than or equal to 10 % of the maximum.

=====										
Nuclide	Te-129m		I-130		Ce-134		Ba-140		La-140	
f1	0.2		1.0		1.0		0.1		0.001	
Class	O		O		D		O		W	

Program	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE
Lungs	2.2E-09	2.2E-09	6.0E-10	5.9E-10	1.2E-08	1.2E-08	1.7E-09	1.6E-09	4.2E-09	4.2E-09
Thyroid			2.0E-08	1.8E-08	1.1E-08	1.1E-08				
Testes										
Ovaries					1.3E-08	1.1E-08	4.3E-10	4.2E-10	4.4E-10	4.5E-10
Red Marrow	8.8E-09	8.6E-09			1.2E-08	1.2E-08	1.3E-09	1.4E-09		
Stomach wall										
SI + contents					1.4E-08	1.4E-08	5.3E-10	5.1E-10	9.7E-10	9.7E-10
ULI wall							1.5E-09	1.4E-09	2.9E-09	2.9E-09
LLI wall	4.2E-09	4.2E-09			1.4E-08	1.4E-08	4.4E-09	4.3E-09	5.5E-09	5.4E-09
Liver										
Kidneys										
Bladder wall										
Muscle					1.1E-08	1.1E-08	2.9E-10	2.8E-10		
Bone Surface	2.0E-08	1.9E-08			1.1E-08	1.1E-08	2.4E-09	2.6E-09		
Skin										
Spleen										
Uterus										
Pancreas										
Total Body										
Remainder					1.5E-08	1.4E-08				

Nuclide	Zr-97		Ce-143		Pr-143		Nd-147		N-239	
f1	0.002		0.0003		0.0003		0.0003		0.01	
Class	Y		Y		O		Y		W	

Program	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE	ICRP	DOSE
Lungs	4.1E-09	4.0E-09	3.9E-09	3.9E-09	1.1E-08	1.1E-08	1.1E-08	1.1E-08	2.4E-09	2.2E-09
Thyroid										
Testes										
Ovaries										
Red Marrow										
Stomach wall										
SI + contents	1.0E-09	1.0E-09								
ULI wall	3.5E-09	3.5E-09	2.1E-09	2.1E-09					1.3E-09	1.2E-09
LLI wall	5.1E-09	5.1E-09	4.3E-09	4.3E-09	6.1E-09	6.1E-09	5.9E-09	5.8E-09	2.9E-09	2.7E-09
Liver										
Kidneys										
Bladder wall										
Muscle										
Bone Surface										
Skin									1.4E-09	1.3E-09
Spleen										
Uterus										
Pancreas										
Total Body										
Remainder										
=====										

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Appendix A
Listing of Computer Programs

ATOMNO_FOR

10-15-1987

```

*****
*
*      SUBROUTINE NAME: ATOMND FORTRAN
*
*      PURPOSE:      Search for atomic number of the nuclide
*
*      INPUT:      Symbol of the radionuclide
*
*****

SUBROUTINE ATOMND (SYM,KZ)
CHARACTER*2 SYM
CHARACTER*2 ZNAME(103)
DATA ZNAME/'H-','HE','LI','BE','B-','C-','N-','O-','F-','NE','NA',
&'MG','AL','SI','P-','S-','CL','AR','K-','CA','SC','TI','V-','CR',
&'MN','FE','CO','NI','CU','ZN','GA','GE','AS','SE','BR','KR','RB','S
&'R','Y-','ZR','NB','MO','TC','RU','RH','PD','AG','CD','IN','SN','SB
&'I','TE','I-','XE','CS','BA','LA','CE','PR','ND','PM','SH','EU','GD'
&'TB','DY','NO','ER','TM','YB','LU','NF','TA','U-','RE','OS','IR',
&'PT','AU','HG','TL','PB','BI','PO','AT','RN','FR','RA','AC','TH',
&'PA','U-','NP','PU','AM','CM','BK','CF','ES','FM','MD','NO','LR'/
*****
*      Comparison
*
*****

DO 5 I=1,103
IF (ZNAME(I) .EQ. SYM) THEN
    KZ=I
    RETURN
END IF
5 CONTINUE
RETURN
END

```

DECAY.FOR

10-15-1987

```

*****
*
*      SUBROUTINE NAME : DECAY FORTRAN
*
*      PURPOSE: Provide the decay scheme of a radionuclide
*
*      DATA FILES REQUIRED:  a) ISOTIPS FILE
*                           b) ISOTOPE FILE
*                           c) ALPHA  FILE
*                           d) BETA  FILE
*                           e) POSITRN FILE
*                           f) ELECTR N FILE
*                           g) PHOTON FILE
*
*****

      SUBROUTINE DECAY (WORD,ICOUNT)
      COMMON EALPHA(1:20),YALPHA(1:20),EBETA(1:50),YBETA(1:50),EPOST(1:
&15),YPOST(1:15),EELEC(1:115),VELEC(1:115),EGAMMA(1:190),YGAMMA(1:
&190),N,I1,I3,I5,I7,NLIFE
      CHARACTER*1 C
      CHARACTER*8 ERT,WORD
      CHARACTER*10 B
      INTEGER DECERR(7)
      OPEN (UNIT=1,FILE='ISOTIPS',ACCESS='DIRECT',RECL=8,FORM='FORMATTED'
&1,STATUS='OLD')
      OPEN (UNIT=2,FILE='ISOTOPE',ACCESS='DIRECT',RECL=59,FORM='FORMATTED'
&2,STATUS='OLD')
      OPEN (UNIT=3,FILE='ALPHA',ACCESS='DIRECT',RECL=20,FORM='FORMATTED'
&3,STATUS='OLD')
      OPEN (UNIT=4,FILE='BETA',ACCESS='DIRECT',RECL=39,FORM='FORMATTED',
&4STATUS='OLD')
      OPEN (UNIT=5,FILE='POSITRN',ACCESS='DIRECT',RECL=31,FORM='FORMATTED'
&5,STATUS='OLD')
      OPEN (UNIT=6,FILE='ELECTRN',ACCESS='DIRECT',RECL=26,FORM='FORMATTED'
&6,STATUS='OLD')
      OPEN (UNIT=7,FILE='PHOTON',ACCESS='DIRECT',RECL=26,FORM='FORMATTED'
&7,STATUS='OLD')
*****
*
*      ISOTIPS
*
*      Comparing the given nuclide name with the alphabetically
*      ordered names (ERT) in the file
*
*****

      ITR=0
      IF (WORD(1:1) .GE. 'R') THEN
        N1=321
      ELSE IF (WORD(1:1) .GE. 'N') THEN
        N1=225
      ELSE IF (WORD(1:1) .GE. 'C') THEN
        N1=65
      ELSE
        N1=1
      END IF
      DO 10 I=N1,496
        READ(1,15,REC=1)ERT

```

DECAY.FOR

10-15-1987

```

15 FORMAT(AB)
   IF (ERT.EQ. WORD) THEN
*****
*
*       ITR-----> Record number of the radionuclide
*
*****
   ITR=1
   GO TO 20
   END IF
10 CONTINUE
   IF (ITR.EQ. 0) THEN
       DECERR(1)=1
       GOTO 80
   END IF
*****
*
*       ISOTOPE
*       DESCRIPTION OF VARIABLES
*       -----
*       ERT-----> Name of the isotope
*       JO-----> Atomic weight
*       J-----> Atomic number
*       B-----> Half-life
*       C-----> Half-life units(S,M,N,D,Y).
*       K-----> Number of daughters
*       L-----> Pointer to first daughter
*       M-----> Number of alphas
*       N-----> Pointer to first alpha
*       I1-----> Number of betas
*       I2-----> Pointer to first beta
*       I3-----> Number of positrons
*       I4-----> Pointer to first positron
*       I5-----> Number of electrons
*       I6-----> Pointer to first electron
*       I7-----> Number of photons
*       I8-----> Pointer to first photon
*
*****
20 READ (2,25,REC=ITR,Iostat=DECERR(2),ERR=80)ERT,JO,J,NLIFE,C,K,L,M,
&N,I1,I2,I3,I4,I5,I6,I7,I8
25 FORMAT(AB,13,I3,G10.0,A1,I1,I3,I2,I3,I2,I4,I2,I3,I3,I4,I3,I4)
*****
*
*       ALPHA
*       AL1-----> Energy in MeV
*       AL2-----> Intensity
*
*****
K1=0
   IF (M.EQ. 0) GOTO 39
   DO 30 JOB=N,N+M-1
       READ(3,35,REC=JOB,Iostat=DECERR(3),ERR=80)AL1,AL2
35 FORMAT(F7.4,1PE13.6)

```

DECAY FOR

10-15-1987

```

K1=K1+1
EALPHA(K1)=AL1
YALPHA(K1)=AL2
30 CONTINUE
*****
*
*           BETA
*
*   BE1-----> Endpoint energy in MeV
*   BE2-----> Average energy in MeV
*   BE3-----> Intensity
*
*****
39 K1=0
IF (I1 .EQ. 0) GOTO 49
DO 40 J1=I2,I2+I1-1
  READ(4,45,REC=J1,IOSTAT=DECERR(4),ERR=80)BE1,BE2,BE3
45 FORMAT(1PE13.6,1PE13.6,1PE13.6)
  K1=K1+1
  EBETA(K1)=BE2
  YBETA(K1)=BE3
40 CONTINUE
*****
*
*           POSITRON
*
*   POS1-----> Endpoint energy in MeV
*   POS2-----> Average energy in MeV
*   POS3-----> Intensity
*
*****
49 K1=0
IF (I3 .EQ. 0) GOTO 59
DO 50 J3=I4,I4+I3-1
  READ(8,55,REC=J3,IOSTAT=DECERR(5),ERR=80)POS1,POS2,POS3
55 FORMAT(F8.5,F10.7,1PE13.6)
  K1=K1+1
  EPOST(K1)=POS2
  YPOST(K1)=POS3
50 CONTINUE
*****
*
*           ELECTRON
*
*   ELE1-----> Energy in MeV
*   ELE2-----> Intensity
*
*****
59 K1=0
IF (I5 .EQ. 0) GOTO 69
DO 60 J4=I6,I6+I5-1
  READ(10,45,REC=J4,IOSTAT=DECERR(6),ERR=80)ELE1,ELE2
65 FORMAT(1PE13.6,1PE13.6)
  K1=K1+1
  EELEC(K1)=ELE1
  YELEC(K1)=ELE2
60 CONTINUE

```

DECAY.FOR

10-15-1987

```
*****
*
*                               PHOTON
*      PH01-----> Energy in MeV
*      PH02-----> Intensity
*
*****

69 K1=0
   IF (I7.EQ. 0)GOTO 79
   DO 70 J5=18,18+I7-1
   READ(11,75,REC=J5,IOSTAT=DECERR(7),ERR=80)PH01,PH02
75  FORMAT(1PE13.6,1PE13.6)
   K1=K1+1
   EGAMMA(K1)=PH01
   YGAMMA(K1)=PH02
70  CONTINUE

*****
*
*      Convert half lives into days
*
*****

79 IF(C.EQ. 'S')THEN
   HLIFE=HLIFE/86400.
ELSE IF (C.EQ. 'M')THEN
   HLIFE=HLIFE/(60.*24.)
ELSE IF (C.EQ. 'H')THEN
   HLIFE=HLIFE/24.
ELSE IF (C.EQ. 'Y')THEN
   HLIFE=HLIFE*365.25
END IF
RETURN
80 CALL CLEAR
   CALL YERROR(DECERR)
   ICOUNT=1
   RETURN
END
```

DECAY1.FOR

10-15-1987

```

*****
*
*   SUBROUTINE NAME: DECAY1 FORTRAN
*   PURPOSE: Find half-lives and names of the given isotope
*             and its daughters
*   DECAY MODE: A---->B---->C---->D
*   DATA FILES REQUIRED:  a) ISOTIPS FILE
*                        b) ISOTOPE FILE
*                        c) DAUTER FILE
*
*   DESCRIPTION OF VARIABLES:
*
*   WORD-----> Name of the given isotope
*   RHALF-----> Vector of half-lives of the given isotope
*                 and its daughters
*   ULIFE-----> Vector of half-life units of the given
*                 isotope and its daughters
*   BRA-----> Vector of branching ratios of the given
*                 isotope (BRA(1)=1) and its daughters
*   RADID-----> Vector of names of the given isotope and
*                 its daughters
*   NO-----> Number of daughters plus one (for the given
*                 isotope
*
*****
      SUBROUTINE DECAY1 (WORD,RHALF,ULIFE,BRA,RADID,NO,*)
      DIMENSION RHALF(1:50),BRA(1:50)
      CHARACTER*1 ULIFE(50),U
      CHARACTER*8 ERT,RADID(50),WORD
      OPEN(UNIT=1,FILE='ISOTIPS',ACCESS='DIRECT',RECL=8,FORM='FORMATTED'
      & )
      OPEN(UNIT=2,FILE='ISOTOPE',ACCESS='DIRECT',RECL=59,FORM='FORMATTED'
      & )
      OPEN(UNIT=12,FILE='DAUTER',ACCESS='DIRECT',RECL=16,FORM='FORMATTED'
      & )
      RADID(1)=WORD
*****
*
*   Data File: ISOTIPS
*
*****
      ITR=0
      IF (WORD(1:1) .GE. 'R') THEN
        M1=321
      ELSE IF (WORD(1:1) .GE. 'H') THEN
        M1=225
      ELSE IF (WORD(1:1) .GE. 'C') THEN
        M1=65
      ELSE
        M1=1
      END IF
      DO 10 I=M1,496
        READ(1,15,REC=1)ERT
15    FORMAT(A8)

```


DECAY1.FOR

10-15-1987

```

IF (ERT .EQ. WORD) THEN
  ITR=1
  GOTD 20
END IF
10 CONTINUE
IF (ITR .EQ. 0) GOTD 50
*****
*
*      Data File: ISOTOPE
*
*****
20 READ(2,25,REC=ITR,ERR=60) ERT,J0,J,B,U,K,L,M,N,I1,I2,I3,I4,I5,I6,I7
  &,I8
*****
*
*      DESCRIPTION OF VARIABLES
*      -----
*
*      ERT---->Name of the isotope
*
*      J0----->Atomic weight
*
*      J----->Atomic number
*
*      B----->Half life
*
*      U----->Half life unite (S,M,H,D,Y)
*
*      K----->Number of daughters
*
*      L----->Pointer to first daughter
*
*      M----->Number of elphas
*
*      N----->Pointer to first elphe
*
*      I1----->Number of betes
*
*      I2----->Pointer to first bete
*
*      I3----->Number of positrons
*
*      I4----->Pointer to first positron
*
*      I5----->Number of electrons
*
*      I6----->Pointer to first electron
*
*      I7----->Number of photons
*
*      I8----->Pointer to first photon
*
*****
25 FORMAT(A8,I3,I3,G10.0,A1,I1,I3,I2,I3,I2,I4,I2,I3,I3,I4,I3,I4)
  RHALF(1)=B
  ULIFE(1)=U
  BRA(1)=1.
*****
*
*      Data File: Dauter
*
*      I9----->Pointer to daughter isotope
*
*      YIELD->Branching ratio of daughter
*
*****
I=1
30 IF (K .EQ. 0) GOTD 45
  YSAVE=0.
  ISAVE=0
*****
*
*      If K > 1 and the decay mode is A----->81, and A----->82,
*

```

DECAY1.FOR

10-15-1987

```

*      then save the pointer and yield of only the higher      *
*      branching ratio                                         *
*
*****
DD 35 J2=L,L-K-1
READ(12,40,REC=J2,ERR=70)I9,YIELD
40 FORMAT(13,1PE13.6)
IF (YIELD .GT. YSAVE)THEN
    YSAVE=YIELD
    ISAVE=I9
END IF
35 CONTINUE
I=I+1
BRA(I)=YSAVE
READ(2,25,REC=ISAVE,ERR=80)ERT,JO,J,B,U,K,L,M,N,11,12,13,14,15,16,
&I7,I8
RADID(I)=ERT
ULIFE(I)=U
RHAF(I)=B
GOTD 30
45 NO=I
RETURN
*****
*
*      ERROR HANDLER
*
*****
50 CALL CLEAR
WRITE (*,55)
55 FORMAT(//,' ERROR: No match found in file "ISOTIPS" for the given
&radionuclide!',' SOURCE: DECAY1 FORTRAN',//,' CORRECTIVE ACTION:
& Try another nuclide!','////)
PAUSE ' TO RESUME PRESS <RETURN>!!'
RETURN 1
60 CALL CLEAR
WRITE (*,65)
65 FORMAT(//,' ERROR: Unable to read the decay scheme from file "ISOT
&OPE" for the given radio nuclide!',' SOURCE: DECAY1 FORTRAN',/
&,' CORRECTIVE ACTION: Try another nuclide!','////)
PAUSE ' TO RESUME PRESS <RETURN>!!'
RETURN 1
70 CALL CLEAR
WRITE (*,75)
75 FORMAT(//,' ERROR: Unable to read the branching ratio and pointer
&of daughter nuclide in file "DAUTER"!',' SOURCE: DECAY1 FORT
&RAN',//,' CORRECTIVE ACTION: Try another nuclide!','////)
PAUSE ' TO RESUME PRESS <RETURN>!!'
RETURN 1
80 CALL CLEAR
WRITE (*,85)
85 FORMAT(//,' ERROR: Unable to read the decay scheme of the daughter
& from file "ISOTOPE"!',' SOURCE: DECAY1 FORTRAN',//,' CORRECTIVE
ACTION: Try another nuclide!','////)
PAUSE ' TO RESUME PRESS <RETURN>!!'

```

DECAY1.FOR

10-15-1987

RETURN 1
END

DOSE.EXE

10-15-1987

FMR SPEFF TEXT
FMR DECAY TEXT
FMR INTRPT TEXT
FMR ENERGY TEXT
FMR FRAC TEXT
FMR DECAY1 TEXT
FMR THALF TEXT
FMR TFRAC TEXT
FMR SOURCE TEXT
FMR SUBMER TEXT
FMR I1 TEXT
FMR TRNSFM TEXT
FMR REFMAN TEXT
FMR INNALE TEXT
FMR RESPIR TEXT
FMR PCLASS TEXT
FMR RESULT TEXT
FMR FACTOR TEXT
FMR INGEST TEXT
FMR YERROR TEXT
FMR DOSE TEXT
FMR ATOMNO TEXT
FMR ICRP TEXT
FMR ICLASS TEXT
FMR F1VALU TEXT
FMR UXP TEXT
EXPAND SUBMER TEXT
EXPAND ATOMNO TEXT
EXPAND SPEFF TEXT
EXPAND DECAY TEXT
EXPAND INTRPT TEXT
EXPAND ENERGY TEXT
EXPAND FRAC TEXT
EXPAND DECAY1 TEXT
EXPAND THALF TEXT
EXPAND TFRAC TEXT
EXPAND SOURCE TEXT
EXPAND I1 TEXT
EXPAND TRNSFM TEXT
EXPAND INNALE TEXT
EXPAND REFMAN TEXT
EXPAND PCLASS TEXT
EXPAND RESPIR TEXT
EXPAND RESULT TEXT
EXPAND FACTOR TEXT
EXPAND INGEST TEXT
EXPAND YERROR TEXT
EXPAND DOSE TEXT
EXPAND ICRP TEXT
EXPAND F1VALU TEXT
EXPAND UXP TEXT
EXPAND ICLASS TEXT
FMR ISOTIPS FILE
FMR ISOTOPE FILE

DOSE.EXE

10-15-1987

FMR ALPHA FILE
FMR BETA FILE
FMR POSITRN FILE
FMR ELECTRN FILE
FMR PHOTON FILE
FMR ABSFRAC FILE
FMR BFFRAC FILE
FMR RETENT FILE
FMR DAUTER FILE
FMR INDEXI FILE
FMR INDEXD FILE
FMR EXCEPT FILE
FMR LIST FILE
FMR NOBLE FILE
FMR CLEAR TEXT
EXPAND CLEAR TEXT
EXPAND ISOTIPS FILE
EXPAND ISOTOPE FILE
EXPAND ALPHA FILE
EXPAND BETA FILE
EXPAND ELECTRN FILE
EXPAND POSITRN FILE
EXPAND PHOTON FILE
EXPAND BFFRAC FILE
EXPAND RETENT FILE
EXPAND DAUTER FILE
EXPAND ABSFRAC FILE
EXPAND INDEXD FILE
EXPAND INDEXI FILE
EXPAND EXCEPT FILE
EXPAND NOBLE FILE
EXPAND LIST FILE
FILEDEF ISOTIPS DISK ISOTIPS FILE A1 (PERM XTENT 496
FILEDEF ISOTOPE DISK ISOTOPE FILE A1 (PERM XTENT 496
FILEDEF ALPHA DISK ALPHA FILE A1 (PERM XTENT 360
FILEDEF BETA DISK BETA FILE A1 (PERM XTENT 1700
FILEDEF POSITRN DISK POSITRN FILE A1 (PERM XTENT 138
FILEDEF ELECTRN DISK ELECTRN FILE A1 (PERM XTENT 3882
FILEDEF PHOTON DISK PHOTON FILE A1 (PERM XTENT 7480
FILEDEF ABSFRAC DISK ABSFRAC FILE A1 (PERM XTENT 4560
FILEDEF BFFRAC DISK BFFRAC FILE A1 (PERM XTENT 501
FILEDEF DAUTER DISK DAUTER FILE A1 (PERM XTENT 291
FILEDEF RETENT DISK RETENT FILE A1 (PERM XTENT 460
FILEDEF INDEXI DISK INDEXI FILE A1 (PERM XTENT 46
FILEDEF INDEXD DISK INDEXD FILE A1 (PERM XTENT 46
FILEDEF EXCEPT DISK EXCEPT FILE A1 (PERM XTENT 695
FILEDEF LIST DISK LIST FILE A1 (PERM XTENT 26
FILEDEF NOBLE DISK NOBLE FILE A1 (PERM XTENT 26
LOAD DOSE
START
ERASE ISOTIPS FILE
ERASE ISOTOPE FILE
ERASE ALPHA FILE
ERASE BETA FILE

DOSE.EXE

10-15-1987

ERASE POSITRN FILE
ERASE ELECTRN FILE
ERASE PHOTON FILE
ERASE DAUTER FILE
ERASE RETENT FILE
ERASE ABSFRAC FILE
ERASE BFACRAC FILE
ERASE SPEFF TEXT
ERASE DECAY TEXT
ERASE INTRPT TEXT
ERASE ENERGY TEXT
ERASE FRAC TEXT
ERASE DECAY1 TEXT
ERASE TNALEF TEXT
ERASE TFRAC TEXT
ERASE SOURCE TEXT
ERASE I1 TEXT
ERASE TRNSFM TEXT
ERASE REFMAN TEXT
ERASE INHALE TEXT
ERASE RESPIR TEXT
ERASE PCLASS TEXT
ERASE RESULT TEXT
ERASE FACTOR TEXT
ERASE CLEAR TEXT
ERASE INGEST TEXT
ERASE YERROR TEXT
ERASE DOSE TEXT
ERASE ATOMND TEXT
ERASE ICLASS TEXT
ERASE FIVALU TEXT
ERASE ICRP TEXT
ERASE LXP TEXT
ERASE EXCEPT FILE
ERASE INDEXI FILE
ERASE INDEXD FILE
ERASE LIST FILE
ERASE NOBLE FILE
ERASE SUBMER TEXT

DOSE.FOR

10-15-1987

```

*****
*
* DESCRIPTION OF VARIABLES:
*
* WORD-----> Name of the given isotope
* KZ -----> Atomic Number of the nuclide
* HFIFTY ----> Specific committed dose equivalent to target
*               organ or tissues
*
*****
      DIMENSION NFIFTY(1:24),MDOSE(1:24),SFACT(19,18),KS(18),KT(19),US(1
&:20,1:50),RHALF(1:50),BRA(1:50),DOSE(1:20)
      INTEGER DOSERR(1:5),OPT,QUI,ORGAN
      CHARACTER*1 SEX,ULIFE(50),CLASS
      CHARACTER*8 WORD,RADID(50),NUCLID,ISOTOP
      CHARACTER*15 UNITS(3)
      CHARACTER*9 SNAME(18)
      CHARACTER*15 TNAME(24)
      CHARACTER*32 UCASE
      CHARACTER*7 FFIL(4)
      5 FORMAT(A)
10  FORMAT(A8,A2,8I2,13,A2,2E10.3,3(A7))
      DATA AMAD,CLASS,F1,KZ,OPT,MAIS,ISAY,MORE,MOM,MU,INRE/1.,'D',0.,0,1
&,2*2,0,2,1,2/
      DATA SEX,QUI/'F',2/
      DATA FFIL /4*'DUMMY' /
      DATA KS/1,2,3,4,5,6,7,8,9,10,11,12,18,13,14,15,16,17/
      DATA KT/12,6,7,8,9,11,10,1,13,4,18,14,5,15,16,3,2,17,19/
      DATA TNAME/'Lungs','Thyroid','Testes','Ovaries','Red Marrow','Stom
&ach wall','SI + contents','ULI wall','LLI wall','Liver','Kidneys','
&Bladder wall','Muscle','Bone Surfaces','Skin','Spleen','Uterus','P
&ancreas','Total Body','Gonads','Adrenals','Lens','Thymus','Brain'
      DATA SNAME/'Bladder','Stomach','SI','ULI','LLI','Kidneys','Liver',
&'Lungs','Muscle','Ovaries','Pancreas','Trab Bone','Skin','Spleen',
&'Testes','Thyroid','Tot. Body','Cort Bone'
      DATA UNITS/'MeV/g','rad/micro Ci.h','mSv/GBq.h'
*****
*
* Display initial screens
*
*****
      CALL CLEAR
      WRITE(*,11)
11  FORMAT(5(/))
      PRINT*, '*****
&*****
      WRITE (*,15)
15  FORMAT(2(/),29X,' D O S E  VERSION 1.0, 1987',/)
      WRITE (*,20)
20  FORMAT(19X,'Written by: Amiruddin Nade')
      WRITE (*,22)
22  FORMAT(19X,'Address:   c/o Dr. R. E. Fau')
      WRITE (*,25)
25  FORMAT(19X,'           Department of Nuclear Engineering')

```

DOSE.FOR

10-15-1987

```

WRITE (*,30)
30 FORMAT(19X,'          Kansas State University')
WRITE (*,35)
35 FORMAT(19X,'          Manhattan, Kansas 66502',//)
PRINT*,/*****
*****/
PAUSE ' TO RESUME PRESS <RETURN>!'
CALL CLEAR
PRINT*,/*****
*****/
WRITE (*,40)
40 FORMAT(//,5X,' PROGRAM NAME: DOSE FORTRAN')
WRITE (*,45)
45 FORMAT(5X,' BASIS:          ICRP Methodology')
WRITE (*,50)
50 FORMAT(5X,' PURPOSE:        TO CALCULATE')
WRITE (*,55)
55 FORMAT(//,8X,'e) Specific committed dose equivalent, HFIFTY (Sv/Bq)
& in target organs')
WRITE (*,60)
60 FORMAT(//,8X,'b) Weighted committed dose equivalent, WDOSE (Sv/Bq)
& in target organs')
WRITE (*,65)
65 FORMAT(//,8X,'c) Annual Limits on Intake, ALI (Bq) of the nuclide')
WRITE (*,70)
70 FORMAT(//,8X,'d) Derived Air Concentration, DAC (Bq/cu.m) of the nu
&clide')
WRITE (*,75)
75 FORMAT(//,8X,'a) Specific Effective Energy Table for 17 sources & 1
&9 targets')
WRITE (*,80)
80 FORMAT(//,8X,'f) Source-organ transformations per unit activity of
&intake',/)
PRINT*,/*****
*****/
85 PAUSE ' TO RESUME PRESS <RETURN>!'
CALL CLEAR
*****
*
*      Additional preliminaries
*
*****
* PRINT *
* PRINT *
* PRINT *, ' 0 - IBM-PC/XT/AT or compatible microcomputer'
* PRINT *, ' 1 - IBM-Mainframe computer, CMS or equivalent system'
* PRINT *
* PRINT *, ' Select integer 0 or 1'
* PRINT *
* READ *, MICRO
*****
110 CALL CLEAR
PRINT *
PRINT *, ' 0 - Date input from the keyboard'

```


DOSE.FOR

10-15-1987

```

PRINT *, ' 1 - Data input from a file'
PRINT *, ' 2 - Preparation of a date input file'
PRINT *
PRINT *, ' Select intagar 0, 1 or 2'
PRINT *
READ (*,115,ERR=110,END=111)NDATA
115 FORMAT(I1)
IF(MDATA .NE. 0 .AND. (NDATA .NE. 1) .AND. (NDATA .NE. 2))THEN
111 REWIND 5
GOTO 110
END IF
IF(MDATA .NE. 0)THEN
120 CALL CLEAR
PRINT *, ' Enter name of date input file (less than or equal to 7
& characters) : '
PRINT *
READ (*,121,ERR=120,END=122)FFILE(3)
121 FORMAT(A7)
IF (FFILE(3) .EQ. ' ')THEN
122 REWIND 5
GOTO 120
END IF
OPEN (75,FILE=FFILE(3),STATUS='UNKNOWN')
REWIND (75)
END IF
*****
*
*      Input date collection
*
*****
IF (NDATA .EQ. 1)GOTO 255
*****
*
*      Identification of nuclide
*
*****
125 CALL CLEAR
PRINT *
PRINT *, 'NOTE: TO INITIATE TERMINATION PRESS <RETURN>!!'
PRINT *
WRITE (*,130)
130 FORMAT(10//), ' Enter radionuclide identification, e.g., CS-137',/)
READ (*,5,ERR=125,END=9000)WORD
NUCLIO=WORD
WORD=UCASE(WORD,1,2)
*****
*
*      Call subroutine OECAY1 for daughters
*
*****
CALL DEAY1 (WORD,RHALF,ULIFE,BRA,RADIO,NO,*125)
OO 162 I=1,NO
ISOTOP=RADIO(I)
*****

```

DOSE.FOR

10-15-1987

```

*          Convert half-lives into days          *
*****
      IF (ULIFE(1) .EQ. 'S') THEN
        RNALF(1)=RNALF(1)/86400.
      ELSE IF (ULIFE(1) .EQ. 'M') THEN
        RNALF(1)=RNALF(1)/(60.*24.)
      ELSE IF (ULIFE(1) .EQ. 'N') THEN
        RNALF(1)=RNALF(1)/24.
      ELSE IF (ULIFE(1) .EQ. 'Y') THEN
        RNALF(1)=RNALF(1)*365.25
      END IF
      IF (I .NE. 1) THEN
        IF (ISOTOP(1:2) .EQ. 'AR' .OR. (ISOTOP(1:2) .EQ. 'KR') .OR. (ISO
&TOP(1:2) .EQ. 'XE') .OR. (ISOTOP(1:2) .EQ. 'NE')) THEN
          NO=1-1
          GOTO 143
        END IF
      END IF
      142 CONTINUE
*****
*          In case of noble radioactive gases or elemental tritium          *
*          the following queries are skipped                                *
*****
      143 IF (WORD(1:2) .EQ. 'H-' .OR. (WORD(1:2) .EQ. 'KR') .OR. (WORD(1:2)
&.EQ. 'XE') .OR. (WORD(1:2) .EQ. 'AR')) THEN
        OPT=3
        GOTO 250
      END IF
*****
*          Identification eex of eubject                                  *
*****
      135 CALL CLEAR
      WRITE (*,140)
      140 FORMAT(10//,' Enter the sex of the exposed individual, e.g., M or
& F',//)
      READ (*,5,ERR=135,END=141)SEX
      IF (SEX .NE.. 'M' .AND. (SEX .NE. 'F')) THEN
        141 REWIND 5
        GOTO 135
      END IF
      SEX=UCASE(SEX,1,1)
*****
*          Query desira for S-matrix table                                *
*****
      145 CALL CLEAR
      WRITE (*,150)
      150 FORMAT(///,' Would you like to eee the specific effective energy t
&able of the nuclide for      17 sources and 19 target organs?',///,

```

DOSE.FOR

10-15-1987

```

& 1 Yea',/, ' 2 No',////, ' Select option by integer---->')
READ (*,115,ERR=145,END=151)OU1
IF (OU1 .NE. 1 .AND. (OU1 .NE. 2))THEN
151 REWIND 5
    GOTD 145
END IF
IF (OU1 .EQ. 1) THEN
155 CALL CLEAR
    PRINT *, ' Enter file specification for table (less than or equal
    & to 7 characters):'
    READ (*,121,ERR=155,END=156)FFILE(1)
    IF (FFILE(1) .EQ. ' ')THEN
156 REWIND 5
        GOTD 155
    END IF
160 CALL CLEAR
    WRITE (*,165)UNITS(1),UNITS(2),UNITS(3)
165 FORMAT(10(//), ' Enter integer for unite selection',/,/, ' 1 ',A,/,/,
    & ' 2 ',A,/,/, ' 3 ',A,/)
    READ (*,115,ERR=160,END=161)NU
    IF (NU .NE. 1 .AND. (NU .NE. 2) .AND. (NU .NE. 3))THEN
161 REWIND 5
        GOTD 160
    END IF
    IF (NU .GT. 1)THEN
170 CALL CLEAR
        PRINT *, ' THE DAUGHTERS OF THE GIVEN NUCLIDE ARE:'
        DO 175 I=2,ND
175 PRINT *, RADID(I)
        PRINT *
        PRINT *
        PRINT *, ' Would you like to see their S-tables too?'
        PRINT *
        PRINT *, ' 1 Yea'
        PRINT *, ' 2 No'
        PRINT *
        PRINT *, ' Select option by integer--->'
        READ (*,115,ERR=170,END=171)MON
        IF (MON .NE. 1 .AND. (MON .NE. 2))THEN
171 REWIND 5
            GOTD 170
        END IF
    END IF
*****
*
*      Query calculation of dose commitments
*
*****
176 CALL CLEAR
    PRINT *, ' Choose one of the followings:'
    PRINT *
    PRINT *, ' 1 Continue data entry for calculation of dose comm
    & itments'
    PRINT *, ' 2 Conclude data entry and STOP'

```

DOSE.FOR

10-15-1987

```

      PRINT *, '      3 Proceed with calculations of S-matrix only'
      IF (NDATA .EQ. 2) THEN
        PRINT *, '      4 Continue data entry for calculation of S-matr
&ix only'
      END IF
      PRINT *
      READ (*,115,ERR=176,END=178) INRE
      IF (INRE .NE. 1 .AND. (INRE .NE. 2) .AND. (INRE .NE. 3) .AND. (I
&NRE .NE. 4)) THEN
178      REWIND 5
          GOTO 176
      END IF
      IF (INRE .EQ. 2) GOTO 9009
      IF (INRE .EQ. 3 .AND. (NDATA .EQ. 0)) GOTO 260
      IF (INRE .NE. 1 .AND. (NDATA .EQ. 2)) THEN
        WRITE(75,10) WORD, SEX, CUI, NU, MON, INRE, OPT, MAIS, ISAY, MORE, KZ, CL
&ASS, AMAD, F1, FFILE(1), FFILE(2), FFILE(4)
        IF (INRE .EQ. 3) THEN
          NDATA=1
          REWIND(75)
        ELSE IF (INRE .EQ. 4) THEN
          GOTO 125
        END IF
      END IF
177  IF (NDATA .EQ. 1) THEN
        READ(75,10,ERR=9009,END=9009) WORD, SEX, CUI, NU, MON, INRE, OPT, MAI
&S, ISAY, MORE, KZ, CLASS, AMAD, F1, FFILE(1), FFILE(2), FFILE(4)
        NUCLID=WORD
        CALL DECAY1 (WORD, RNALF, ULIFE, BRA, RADIO, NO, *9009)
        DO 179 I=1, NO
          ISOTOP=RADIO(I)
*****
*      Convert half-lives into days      *
*****
          IF (ULIFE(I) .EQ. 'S') THEN
            RNALF(I)=RNALF(I)/86400.
          ELSE IF (ULIFE(I) .EQ. 'M') THEN
            RNALF(I)=RNALF(I)/(60.*24.)
          ELSE IF (ULIFE(I) .EQ. 'Y') THEN
            RNALF(I)=RNALF(I)/24.
          ELSE IF (ULIFE(I) .EQ. 'Y') THEN
            RNALF(I)=RNALF(I)*365.25
          END IF
          IF (I .NE. 1) THEN
            IF (ISOTOP(1:2) .EQ. 'AR' .OR. (ISOTOP(1:2) .EQ. 'KR') .OR. (ISO
&TOP(1:2) .EQ. 'XE') .OR. (ISOTOP(1:2) .EQ. 'NE')) THEN
              NO=I-1
              GOTO 260
            END IF
          END IF
179      CONTINUE
          GOTO 260
        END IF
      END IF

```

DOSE.FOR

10-15-1987

```

*****
*
*      Identification of mode of intake
*
*****

180 CALL CLEAR
    WRITE(*,185)
185 FORMAT(10(//),' MODE OF INTAKE OF THE RADIONUCLIDE:',/)
    WRITE (*,190)
190 FORMAT(/,8X,'1. ....Ingestion')
    WRITE (*,195)
195 FORMAT(8X,'2. ....Inhalation')
    WRITE (*,205)
205 FORMAT(///,' Select Option by Integer ----->',/)
    READ (*,115,ERR=180,END=181)OPT
    IF (OPT .LT. 1 .OR. (OPT .GT. 2))THEN
181      REWIND 5
      GOTO 180
    END IF
    IF (WORD(1:2) .EQ. 'AR' .OR. (WORD(1:2) .EQ. 'KR') .OR. (WORD(1:
&2) .EQ. 'XE') .OR. (WORD(1:2) .EQ. 'N-'))THEN
      DOSERR(2)=1
      GOTO 1000
    END IF
210 CALL CLEAR
    WRITE (*,215)
215 FORMAT(5(//),' Would you like to see the number of transformation
&s of the nuclide in source organs? ',/,,' 1 Yes',/,,' 2 N
&o',/,/,,' Select option by Integer --->',/,/)
    READ (*,115,ERR=210,END=211)MAIS
    IF (MAIS .NE. 1 .AND. (MAIS .NE. 2))THEN
211      REWIND 5
      GOTO 210
    END IF
    IF (MAIS .EQ. 1)THEN
216      CALL CLEAR
      PRINT *, ' Enter file specification for transformations: '
      PRINT *, ' (less than or equal to 7 characters !)'
      READ (*,121,ERR=216,END=217)FFILE(4)
      IF (FFILE(4) .EQ. ' ')THEN
217        REWIND 5
          GOTO 216
      END IF
      IF (NO .GT. 1)THEN
220        CALL CLEAR
          WRITE (*,225)
225        FORMAT(5(//),' Would you like to see the number of transfor
&ations of the daughters too? ',/,,' 1 Yes',/,,' 2 No',/,/,/,
&' SELECT OPTION BY INTEGER--->',/,/)
          READ (*,115,ERR=220,END=221)ISAY
          IF (ISAY .NE. 1 .AND. (ISAY .NE. 2))THEN
221            REWIND 5
              GOTO 220
          END IF

```

DOSE.FOR

10-15-1987

```

      END IF
      END IF
*****
*
*      Atomic Number of the given radionuclide
*
*****
      KZ=0
      CALL ATOMNO (WORD(1:2),KZ)
      IF (KZ .EQ. 0)THEN
        DOSERR(3)=1
        GOTO 1000
      END IF
*****
*
*      Obtain data for inhalation class and transfer to body fluid
*
*****
      IF (OPT .EQ. 2)THEN
        CALL ICLASS(CLASS)
      END IF
      CALL F1VALU(KZ,F1,*125)
*****
*
*      Identification of aerodynamic diameter for inhalation
*
*****
      IF (OPT .EQ. 2)THEN
230  CALL CLEAR
        WRITE (*,235)
235  FORMAT(//,' Is the activity median aerodynamic diameter equal to
        & 1 micrometer?',//,' 1 Yes',//,' 2 No',////,' Select option by inte
        & ger--->',/)
        READ (*,115,ERR=230,END=231)LOT
        IF (LOT .NE. 1 .AND. (LOT .NE. 2))THEN
231  REWIND 5
          GOTO 230
        END IF
        IF (LOT .EQ. 2)THEN
240  CALL CLEAR
          WRITE (*,245)
245  FORMAT(/,' Enter the value of AMAD (micrometers) between 0.1
        & end 20 micrometers !')
          READ(*,ERR=240,END=241)AMAD
          IF (AMAD .LT. 0.1 .OR. (AMAD .GT. 20.))THEN
241  REWIND 5
            GOTO 240
          END IF
        END IF
      END IF
*****
*
*      Query filename for output table
*
*****

```

DOSE.FOR

10-15-1987

```

*****
250 CALL CLEAR
   PRINT *, ' Enter file specification for dose commitment results:'
   PRINT *, ' (less than or equal to 7 characters )'
   READ (*,121,ERR=250,END=252)FFILE(2)
   IF (FFILE(2) .EQ. ' ') THEN
252   REWIND 5
      GOTD 250
   END IF
   IF (NDATA .NE. 2) THEN
      IF (WORD(1:2) .EQ. 'AR' .OR. (WORD(1:2) .EQ. 'KR') .OR. (WORD(1:
&2) .EQ. 'XE') .OR. (WORD(1:2) .EQ. 'H-')) THEN
         GOTD 268
      END IF
*****
*
*      Query continuation of data input
*
*****
ELSE
251 CALL CLEAR
   PRINT *
   PRINT *, ' 0 - Conclude data entry and STOP'
   PRINT *, ' 1 - Continue data entry'
   PRINT *, ' 2 - Proceed with calculations'
   PRINT *
   PRINT *, ' Select integer 0, 1, or 2'
   PRINT *
   READ (*,115,ERR=251,END=253)MORE
   IF (MORE .NE. 0 .AND. (MORE .NE. 1) .AND. (MORE .NE. 2)) THEN
253   REWIND 5
      GOTD 251
   END IF
   WRITE(75,10)WORD,SEX,OUI,MU,MON,INRE,OPT,MAIS,ISAT,MORE,KZ,CLASS
&,AMAD,F1,FFILE(1),FFILE(2),FFILE(4)
   IF (MORE .EQ. 1) THEN
      GOTD 125
   ELSE IF (MORE .EQ. 2) THEN
      NDATA=1
      REWIND(75)
   ELSE
      CALL CLEAR
      GOTD 9009
   END IF
   END IF
255 IF (NDATA .EQ. 1) THEN
   READ(75,10,ERR=9009,END=9009)WORD,SEX,OUI,MU,MON,INRE,OPT,MAIS,I
&SAT,MORE,KZ,CLASS,AMAD,F1,FFILE(1),FFILE(2),FFILE(4)
   NUCLID=WORD
   IF (OPT .EQ. 3) THEN
      IF (WORD(1:2) .NE. 'AR' .AND. (WORD(1:2) .NE. 'KR') .AND. (WORD(
&1:2) .NE. 'XE') .AND. (WORD(1:2) .NE. 'H-')) THEN
         DOSERR(2)=1
         GOTD 1000

```

DOSE.FOR

10-15-1987

```

ELSE
  GOTO 268
END IF
ELSE IF (OPT .EQ. 3) THEN
  IF (WORD(1:2) .EQ. 'AR' .OR. (WORD(1:2) .EQ. 'KR') .OR. (WORD(1:
&2) .EQ. 'XE') .OR. (WORD(1:2) .EQ. 'H-')) THEN
    DOSERR(2)=1
    GOTO 1000
  END IF
  CALL DECAY1 (WORD,RHALF,ULIFE,BRA,RADIO,NO,"9009")
  DO 256 I=1,NO
    ISOTOP=RADIO(I)
*****
*      Convert half-lives into days      *
*****
    IF (ULIFE(I) .EQ. 'S') THEN
      RHALF(I)=RHALF(I)/86400.
    ELSE IF (ULIFE(I) .EQ. 'M') THEN
      RHALF(I)=RHALF(I)/(60.*24.)
    ELSE IF (ULIFE(I) .EQ. 'H') THEN
      RHALF(I)=RHALF(I)/24.
    ELSE IF (ULIFE(I) .EQ. 'Y') THEN
      RHALF(I)=RHALF(I)*365.25
  END IF
  IF (I .EQ. 1) THEN
    IF (ISOTOP(1:2) .EQ. 'AR' .OR. (ISOTOP(1:2) .EQ. 'KR') .OR. (ISO
&TOP(1:2) .EQ. 'XE') .OR. (ISOTOP(1:2) .EQ. 'H-')) THEN
      NO=I-1
      GOTO 260
    END IF
  END IF
  ENO IF
256 CONTINUE
END IF
END IF
*****
*
*      Calculation and printing results of S-matrix      *
*****
260 CALL CLEAR
  IF (OUT .EQ. 1) THEN
    IF (MON .EQ. 2) THEN
      MES=1
    ELSE
      MES=NO
    END IF
  END IF
  OPEN(55,FILE=FFILE(1),STATUS='UNKNOWN')
  DO 265 I=1,MES
    PRINT *, ' Calculating S-matrix for ',RADIO(I)
    WRITE (*,270)FFILE(1)
270  FORMAT(//,' Results are in file ',A)
    CALL FACTOR (RADIO(I),NU,SFACT,NOATA,WORD,RHALF(1),*125,"9009")
    CALL CLEAR
    WRITE (55,275)

```


DOSE.FOR

10-15-1987

```

      IF (NDATA .EQ. 0) THEN
        WRITE (*,275)
      END IF
275  FORMAT(/, ' *****
      &*****',/,6X,'*',64K,'*',/,6X,'*' S-FACTORS FOR THE AD
      &ULT BY METHODS OF ICRP-30, V. 1.0, 1987  ',/,6X,'*',64K,'*',/,6X
      &,'*' Based on : "Radioactive Decay Data Tables",18X,'*',/,6X,'*'
      &,'16X,'O.C. Kocher, DOE/TIC-11026 (1981)',15X,'*',/,6X,'*' Written
      & by : Amiruddin Nade',34X,'*',/,6X,'*' Address : c/o Dr. R.E.
      & Fah',32X,'*',/,6X,'*',16X,'Department of Nuclear Engineering',15K
      &,'*',/,6X,'*',16X,'Kansas State University',25K,'*',/,6X,'*',16X,'
      &Manhattan, Kansas 66506',25K,'*',/,6X,'*',64K,'*',/, ' *****
      &*****')
      WRITE (55,280)RADIO(I),UNITS(MU)
      IF (NDATA .EQ. 0) THEN
        WRITE (*,280) RADIO(I),UNITS(MU)
      END IF
280  FORMAT(/,21K,A,' S-FACTORS (' ,A,')')
      WRITE (55,285)
      IF (NDATA .EQ. 0) THEN
        WRITE (*,285)
      END IF
285  FORMAT(/,9X,'TARGET',28X,'SOURCE ORGANS')
      WRITE (55,290)(SNAME(KS(J)),J=1,4)
      IF (NDATA .EQ. 0) THEN
        WRITE (*,290)(SNAME(KS(J)),J=1,4)
      END IF
290  FORMAT(9X,'ORGAN',11K,4(3K,A),/)
      DO 295 K=1,19
        WRITE (55,300) TNAME(KT(K)),(SFAC(T(K),KS(J)),J=1,4)
        IF (NDATA .EQ. 0) THEN
          WRITE (*,300)TNAME(KT(K)),(SFAC(T(K),KS(J)),J=1,4)
        END IF
295  CONTINUE
300  FORMAT(9X,A,4(3K,1PE9.2))
      WRITE (55,305)
      IF (NDATA .EQ. 0) THEN
        WRITE (*,305)
      END IF
305  FORMAT(1H1)
      WRITE (55,280)RADIO(I),UNITS(MU)
      WRITE (55,285)
      WRITE (55,290)(SNAME(KS(J)),J=5,8)
      IF (NDATA .EQ. 0) THEN
        WRITE (*,280) RADIO(I),UNITS(MU)
        WRITE (*,285)
        WRITE (*,290)(SNAME(KS(J)),J=5,8)
      END IF
      DO 310 K=1,19
        WRITE (55,300) TNAME(KT(K)),(SFAC(T(K),KS(J)),J=5,8)
        IF (NDATA .EQ. 0) THEN
          WRITE (*,300) TNAME(KT(K)),(SFAC(T(K),KS(J)),J=5,8)
        END IF
310  CONTINUE

```

DOSE. FOR

10-15-1987

```
WRITE (55,305)
WRITE (55,280)RADID(1),UNITS(MU)
WRITE (55,285)
WRITE (55,290)(SNAME(KS(J)),J=9,12)
IF (NDATA .EQ. 0)THEN
  WRITE (*,305)
  WRITE (*,280) RADID(1),UNITS(MU)
  WRITE (*,285)
  WRITE (*,290)(SNAME(KS(J)),J=9,12)
END IF
DO 315 K=1,19
  WRITE (55,300) TNAME(KT(K)),(SFACT(KT(K),KS(J)),J=9,12)
  IF (NDATA .EQ. 0)THEN
    WRITE (*,300)TNAME(KT(K)),(SFACT(KT(K),KS(J)),J=9,12)
  END IF
315 CONTINUE
  WRITE (55,305)
  WRITE (55,280)RADID(1),UNITS(MU)
  WRITE (55,285)
  WRITE (55,290)(SNAME(KS(J)),J=13,16)
  IF (NDATA .EQ. 0)THEN
    WRITE (*,305)
    WRITE (*,280) RADID(1),UNITS(MU)
    WRITE (*,285)
    WRITE (*,290)(SNAME(KS(J)),J=13,16)
  END IF
  DO 320 K=1,19
    WRITE (55,300) TNAME(KT(K)),(SFACT(KT(K),KS(J)),J=13,16)
    IF (NDATA .EQ. 0)THEN
      WRITE (*,300) TNAME(KT(K)),(SFACT(KT(K),KS(J)),J=13,16)
    ENDIF
320 CONTINUE
    WRITE (55,305)
    WRITE (55,280)RADID(1),UNITS(MU)
    WRITE (55,321)
    WRITE (55,322)(SNAME(KS(J)),J=17,18)
    IF (NDATA .EQ. 0)THEN
      WRITE (*,305)
      WRITE (*,280) RADID(1),UNITS(MU)
      WRITE (*,321)
      WRITE (*,322)(SNAME(KS(J)),J=17,18)
    END IF
321 FORMAT(/,9X,'TARGET',14X,'SOURCE ORGANS')
322 FORMAT(9X,'ORGAN',11X,2(3X,A),/)
    DO 323 K=1,19
      WRITE (55,300) TNAME(KT(K)),(SFACT(KT(K),KS(J)),J=17,18)
      IF (NDATA .EQ. 0)THEN
        WRITE (*,300) TNAME(KT(K)),(SFACT(KT(K),KS(J)),J=17,18)
      END IF
323 CONTINUE
    WRITE (55,305)
    IF (NDATA .EQ. 0)THEN
      WRITE (*,305)
    END IF
```

DOSE.FOR

10-15-1987

```

265  CONTINUE
      END IF
      IF (INRE .EQ. 4)GOTO 177
      IF (INRE .EQ. 3 .AND. (NDATA .EQ. 0))GOTO 125
      IF (INRE .EQ. 3 .AND. (NDATA .NE. 0))GOTO 9009
*****
*                                     *
*      Calculation of Dose Commitments      *
*                                     *
*****
      IF (OPT .EQ. 3)THEN
        IF (WORD(1:2) .NE. 'AR' .OR. (WORD(1:2) .NE. 'KR') .OR. (WORD(1:
&2) .NE. 'NE') .OR. (WORD(1:2) .NE. 'N-')) THEN
          DOSERR(2)=1
          GOTO 1000
        END IF
      ELSE
        IF (WORD(1:2) .EQ. 'AR' .OR. (WORD(1:2) .EQ. 'KR') .OR. (WORD(1:
&2) .EQ. 'NE') .OR. (WORD(1:2) .EQ. 'N-'))THEN
          DOSERR(2)=1
          GOTO 1000
        END IF
      END IF
      END IF
268  CALL CLEAR
      IF (OPT .EQ. 1)THEN
        PRINT *, ' Calculating ingestion dose for ',NUCLID
        PRINT *, ' F1 (GI to body fluids) = ',F1
        PRINT *, ' Subject = ',SEX
      ELSE IF (OPT .EQ. 2)THEN
        PRINT *, ' Calculating inhalation dose for ',NUCLID
        PRINT *, ' Inhalation class = ',CLASS
        PRINT *, ' F1 (GI to body fluids) = ',F1
        PRINT *, ' Subject = ',SEX
      ELSE IF (OPT .EQ. 3)THEN
        PRINT *, ' Calculating submerison dose for ',NUCLID
      END IF
      DER=D.
      RISK=0.
      ORGAN=0
      CALL ICRP(OPT,WORD,SEX,F1,CLASS,AMAD,K2,NFIFTY,US,ROB,NDATA,DER,R1
&SK,ORGAN,*125,*9009)
      IF (WORD(1:2) .EQ. 'AR' .OR. (WORD(1:2) .EQ. 'KR') .OR. (WORD(1:
&2) .EQ. 'XE') .OR. (WORD(1:2) .EQ. 'N-'))THEN
        GOTO 366
      END IF
      IF (NUCLID(1:2) .EQ. 'BA' .OR. (NUCLID(1:2) .EQ. 'RA') .OR. (NUCLID
&(1:2) .EQ. 'SR') .OR. (NUCLID(1:2) .EQ. 'CA'))THEN
        ROB=65000.
      ELSE IF (NUCLID(1:2) .EQ. 'RE' .OR. (NUCLID(1:2) .EQ. 'TC'))THEN
        ROB=68030.
      ELSE IF (NUCLID(1:2) .EQ. 'C-')THEN
        ROB=70000.
      ELSE IF (NUCLID .EQ. 'TE-131M ' .OR. (NUCLID .EQ. 'TE-131 ' ) .OR.
& (NUCLID .EQ. 'TE-132 ' ) .OR. (NUCLID .EQ. 'TE-133 ' ) .OR. (NUCLID

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DOSE.FOR

10-15-1987

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&ID .EQ. 'TE-133 ') .OR. (NUCLID .EQ. 'TE-133M ') .OR. (NUCLID .EQ.
& 'TE-134 ') THEN
  ROB=64980.
  END IF
*****
*
*   Print results of source-organ transformations
*
*****
325 CALL CLEAR
  IF (MAIS .EQ. 1) TXEH
  IF (ISAY .EQ. 2) TXEH
  MES=1
  ELSE
  MES=NO
  END IF
  OPEX(85, FILE=FFILE(4), STATUS='UNKNOWN')
  WRITE (*, 335) FFILE(4)
335  FORMAT(//, ' Results of source transformations are in file ', A)
  DO 330 I=1, MES
  ISOTOP=RADIO(I)
  IF (ULIFE(I) .EQ. 'S') TXEH
    RHALF(I)=RHALF(I)/86400.
  ELSE IF (ULIFE(I) .EQ. 'H') THEN
    RHALF(I)=RHALF(I)/(60.*24.)
  ELSE IF (ULIFE(I) .EQ. 'X') TXEH
    RHALF(I)=RHALF(I)/24.
  ELSE IF (ULIFE(I) .EQ. 'Y') TXEH
    RHALF(I)=RHALF(I)*365.25
  END IF
  WRITE (85, 340) RADIO(I)
  IF (XDATA .EQ. 0) TXEH
  WRITE (*, 340) RADIO(I)
  END IF
340  FORMAT(//, ' *****
&*****', /, 6X, ' ', 6X, ' ', 6X, ' ' SOURCE-ORGAN TRA
&SFORMATIONS OF ', A, ' V. 1.0, 1987', 6X, ' ', 6X, ' ', 6X, ' ', 6X, ' '
& Based on : "Radioactive Decay Data Tables", 18X, ' ', 6X, ' ', 1
&6X, 'D.C. Kocher, DOE/TIC-11026 (1981)', 15X, ' ', 6X, ' ' Written by
& : Amiruddin Xuda', 34X, ' ', 6X, ' ' Address : c/o Dr. R.E. F
&u', 32X, ' ', 6X, ' ', 16X, 'Department of Nuclear Engineering', 15X, ' '
&', /, 6X, ' ', 16X, 'Kansas State University', 25X, ' ', 6X, ' ', 16X, 'Man
&hattan, Kansas 66506', 25X, ' ', 6X, ' ', 6X, ' ', 6X, ' ', 6X, ' ', 6X, ' '
&*****', //)
  IF (XDATA .EQ. 0) TXEH
  WRITE (*, 375) RADIO(I)
  END IF
  WRITE (85, 375) RADIO(I)
  IF (SEX .EQ. 'M') THEN
  IF (NDATA .EQ. 0) TXEH
  WRITE (*, 380)
  END IF
  WRITE (85, 380)
  ELSE

```

DOSE.FOR

10-15-1987

```

WRITE (85,385)
IF (NDATA .EQ. 0) THEN
  WRITE (*,385)
END IF
END IF
IF (OPT .EQ. 2) THEN
  WRITE(85,388)
  WRITE(85,390) CLASS
  WRITE(85,395) AMAD
  WRITE(85,400) F1
  IF (NDATA .EQ. 0) THEN
    WRITE(*,388)
    WRITE(*,390) CLASS
    WRITE(*,395) AMAD
    WRITE(*,400) F1
  END IF
ELSE IF (OPT .EQ. 1) THEN
  WRITE(85,420)
  WRITE(85,425) F1
  IF (NDATA .EQ. 0) THEN
    WRITE(*,420)
    WRITE(*,425) F1
  END IF
END IF
WRITE (85,345)
IF (NDATA .EQ. 0) THEN
  WRITE (*,345)
END IF
345 FORMAT(////,20X,' SOURCE ORGAN',10X,' TRANSFORMATIONS (/Bq)',/)
IF (US(12,1) .EQ. 0.) GOTD 350
*****
*   Nuclides uniformly distributed in volume of mineral bone   *
*****
  IF (NUCLID(1:4) .EQ. 'P-33' .OR. (NUCLID(1:6) .EQ. 'NB-93M') .OR.
& (NUCLID(1:5) .EQ. 'NB-94') .OR. (NUCLID(1:5) .EQ. 'U-232') .OR.
& (NUCLID(1:5) .EQ. 'U-233') .OR. (NUCLID(1:5) .EQ. 'U-234') .OR. (NUCLID(1:5) .EQ. 'U-235') .OR. (NUCLID(1:5) .EQ. 'U-236') .OR. (NUCLID(1:5) .EQ. 'U-238') .OR. (NUCLID(1:2) .EQ. 'NA') .OR. (NUCLID(1:2) .EQ. 'CR') .OR. (NUCLID(1:2) .EQ. 'RB') .OR. (NUCLID(1:5) .EQ. 'Zn-65') .OR. (NUCLID(1:6) .EQ. 'PB-205') .OR. (NUCLID(1:6) .EQ. 'PB-210') .OR. (NUCLID(1:4) .EQ. 'BE-7') .OR. (NUCLID(1:5) .EQ. 'B&E-10') .OR. (NUCLID(1:4) .EQ. 'V-49') .OR. (NUCLID(1:6) .EQ. 'PD-1&03') .OR. (NUCLID(1:6) .EQ. 'PD-107') .OR. (NUCLID(1:6) .EQ. 'SN-1&13') .OR. (NUCLID(1:7) .EQ. 'SN-119M') .OR. (NUCLID(1:6) .EQ. 'SN-1&123') .OR. (NUCLID(1:6) .EQ. 'SN-126') .OR. (NUCLID(1:6) .EQ. 'TA-1&82') .OR. (NUCLID(1:5) .EQ. 'W-181') .OR. (NUCLID(1:5) .EQ. 'W-18&5') .OR. (NUCLID(1:5) .EQ. 'W-188')) THEN
    US(12,1) = US(12,1)*0.2
    US(18,1) = (US(12,1)/0.2)*0.8
*****
*   Alkaline earths   *
*****
  ELSE IF (NUCLID(1:2) .EQ. 'SR' .OR. (NUCLID(1:2) .EQ. 'BA') .OR. (
& NUCLID(1:2) .EQ. 'CA') .OR. (NUCLID(1:2) .EQ. 'RA')) THEN

```

DOSE.FOR

10-15-1987

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      IF (RHALF(1) .GT. 15) THEN
        US(12,1)=US(12,1)*0.2
        US(18,1)=(US(12,1)/0.2)*0.8
      ELSE
        US(12,1)=US(12,1)*0.5
        US(18,1)=US(12,1)
      END IF
    ELSE
      US(12,1)=US(12,1)*0.5
      US(18,1)=US(12,1)
    END IF
350   GO 355 K=1,18
      IF (US(K,1) .EQ. 0.) GO 355
      IF (K .EQ. 17) THEN
        IF (ROB .LT. 70000.) THEN
          WRITE(85,351)US(K,1)*86400.
          IF (NDATA .EQ. 0) THEN
            WRITE(*,351)US(K,1)*86400.
          END IF
351   FORMAT(23X,'Other tissue',11X,1PE9.2)
          GO 355
        END IF
      END IF
      WRITE(85,360)SNAME(K),US(K,1)*86400.
      IF (NDATA .EQ. 0) THEN
        WRITE(*,360)SNAME(K),US(K,1)*86400.
      END IF
360   FORMAT(23X,A,14X,1PE9.2)
355   CONTINUE
362   FORMAT(//,16X,' Mass of other tissue = ',F9.2,' grams')
      IF (ROB .LT. 70000.) THEN
        IF (NDATA .EQ. 0) THEN
          WRITE(*,362)ROB
        END IF
      END IF
      WRITE (85,305)
      IF (NDATA .EQ. 0) THEN
        WRITE (*,305)
      END IF
330   CONTINUE
      END IF
*****
*
*       Print results of dose commitments
*
*****
      IF (NOATA .EQ. 0) THEN
        PAUSE 'TO RESUME PRESS <RETURN>|'
      END IF
366   CALL CLEAR
      OPEN (65,FILE=FFILE(2),STATUS='UNKNOWN')
      WRITE (*,365)FFILE(2)
365   FORMAT(//,' Results of Dose Commitments are in file ',A)

```


DOSE.FOR

10-15-1987

```

ELSE IF (I .EQ. 20) THEN
  WT=0.25
ELSE IF (I .EQ. 5) THEN
  WT=0.12
ELSE IF (I .EQ. 13) THEN
  WT=0.15
ELSE IF (I .EQ. 14) THEN
  WT=0.03
ELSE
  WT=0.06
END IF
IF (I .EQ. 15 .OR. (I .EQ. 22)) THEN
  IF (NDATA .EQ. 0) THEN
    WRITE(*,443) TNAME(I), NFIFTY(I)
  END IF
  WRITE(65,443) TNAME(I), NFIFTY(I)
443 FORMAT(6X,' ',2X,A14,6X,1PEB.1,7X,' ',26X,' ')
ELSE
  IF (NDATA .EQ. 0) THEN
    WRITE(*,444) TNAME(I), NFIFTY(I), WT*NFIFTY(I)
  END IF
  WRITE(65,444) TNAME(I), NFIFTY(I), WT*NFIFTY(I)
444 FORMAT(6X,' ',2X,A14,6X,1PEB.1,7X,' ',9X,1PEB.1,9X,' ')
END IF
442 CONTINUE
IF (RISK .EQ. 0) THEN
  IF (NDATA .EQ. 0) THEN
    WRITE(*,446) DER
  END IF
  WRITE(65,446) DER
446 FORMAT(6X,' ',37X,' ',26X,' ',/,6X,'*****
&*****',/,6X,' ',64X,' ',/,6X,' '
&5X,'Stochastic Risk',8X,'Non-Stochastic Risk (Drgen)',9X,' ',/,6
&X,' ',6X,1PEB.1,' Bq/m^3 ',10X,1PEB.1,' Bq/m^3 ',1X,' (' ,A14,' )',1X
&,' ',/,6X,' ',64X,' ',/,6X,'*****
&*****')
ELSE
  IF (NDATA .EQ. 0) THEN
    WRITE(*,447) DER, RISK, TNAME(ORGAN)
  END IF
  WRITE(65,447) DER, RISK, TNAME(ORGAN)
447 FORMAT(6X,' ',37X,' ',26X,' ',/,6X,'*****
&*****',/,6X,' ',64X,' ',/,6X,' '
&5X,'Stochastic Risk',8X,'Non-Stochastic Risk (Drgen)',9X,' ',/,6
&X,' ',6X,1PEB.1,' Bq/m^3 ',10X,1PEB.1,' Bq/m^3 ',1X,' (' ,A14,' )',1X
&,' ',/,6X,' ',64X,' ',/,6X,'*****
&*****')
END IF
GOTO 480
END IF
IF (SEX .EQ. 'M') THEN
  IF (NDATA .EQ. 0) THEN
    WRITE(*,380)
  END IF
  WRITE(65,380)

```


DOSE.FOR

10-15-1987

```
380 FORMAT(10X,'Subject : ',22X,'Male')
      ELSE
        WRITE (65,385)
        IF (NDATA .EQ. D)THEN
          WRITE (*,385)
        END IF
385  FORMAT(10X,'Subject : ',22X,'Female')
      END IF
      IF (OPT .EQ. 2)THEN
        WRITE(65,388)
        IF (NDATA .EQ. 0)THEN
          WRITE (*,388)
        END IF
388  FORMAT(10X,'Mode of intake : ',15X,'Inhalation')
        WRITE(65,390)CLASS
        IF (NDATA .EQ. 0)THEN
          WRITE(*,390)CLASS
        END IF
390  FORMAT(10X,'Inhalation Class : ',13X,A)
        WRITE(65,395)AMAD
        IF (NDATA .EQ. D)THEN
          WRITE(*,395)AMAD
        END IF
395  FORMAT(10X,'Particle AMAD (um) : ',10X,F5.2)
        WRITE(65,400)F1
        IF (NDATA .EQ. D)THEN
          WRITE(*,400)F1
        END IF
400  FORMAT(10X,'Body fluid transfer fraction : ',F8.5)
        IF (ND .GT. 1 .AND. (ND .LT. 4))THEN
          WRITE(65,405)(RADID(I),I=2,NO)
          WRITE(65,415)(BRA(I),I=2,NO)
          IF (NDATA .EQ. 0)THEN
            WRITE(*,405)(RADID(I),I=2,NO)
            WRITE(*,415)(BRA(I),I=2,NO)
          END IF
405  FORMAT(10X,'Daughter Products : ',12X,3A)
415  FORMAT(10X,'Branching Ratios : ',12X,3(F6.3,2X))
      ELSE IF (NO .GE. 4)THEN
        WRITE (65,406)
        IF (NDATA .EQ. 0)THEN
          WRITE (*,406)
        END IF
        DO 407 I=2,NO
          WRITE (65,408)RADID(I),BRA(I)
          IF (NDATA .EQ. D)THEN
            WRITE (*,408)RADID(I),BRA(I)
          END IF
407  CONTINUE
        WRITE (65,305)
        IF (NDATA .EQ. D)THEN
          WRITE (*,305)
        END IF
406  FORMAT(10X,'Daughter Products',12X,'Branching Ratios')
```

DOSE.FOR

10-15-1987

```

408   FORMAT(14X,A,21X,F6.3)
      ENO IF
      ELSE IF (OPT .EQ. 1) THEN
        WRITE(65,420)
        WRITE(65,425)F1
        IF (NDATA .EQ. 0) THEN
          WRITE(*,420)
          WRITE(*,425)F1
420   FORMAT(10X,'Mode of intake : ',15X,'Ingestion')
425   FORMAT(10X,'Body fluid transfer fraction : ',F8.5)
      END IF
      IF (NO .GT. 1 .AND. (NO .LT. 4)) THEN
        WRITE(65,426)(RADIO(I),I=2,NO)
        WRITE(65,428)(BRA(I),I=2,NO)
        IF (NDATA .EQ. 0) THEN
          WRITE(*,426)(RADIO(I),I=2,NO)
          WRITE(*,428)(BRA(I),I=2,NO)
        END IF
426   FORMAT(10X,'Daughter Products : ',12X,3A)
428   FORMAT(10X,'Branching Ratios : ',12X,3(F6.3,2X))
      ELSE IF (NO .GE. 4) THEN
        WRITE (65,406)
        IF (NDATA .EQ. 0) THEN
          WRITE (*,406)
          ENO IF
          DO 429 I=2,NO
            WRITE (65,408)RADIO(I),BRA(I)
            IF (NDATA .EQ. 0) THEN
              WRITE (*,408)RADIO(I),BRA(I)
            END IF
429   CONTINUE
            WRITE (65,305)
            IF (NDATA .EQ. 0) THEN
              WRITE (*,305)
            END IF
          ENO IF
        END IF
        DO 430 I=1,20
          DOSE(I)=HFIFTY(I)
430   CONTINUE
          CALL RESULT(NFIFTY,DOSE,ALI,POST,IRGANT,OAC,KZ,REMDR,WREMDR,WT
            &F,SUM)
          IF (NDATA .EQ. 0) THEN
            WRITE(*,440)
            ENO IF
            WRITE(65,440)
440   FORMAT(/,
            &*****
            &' ,6X,' ',2X,'Target',6X,'Specific Committed', ' '
            &' , ' Ooses greater than or equal  ',6X,' ',2X,'Organ ',6X,'Oose
            &Equivalent',4X,' ', ' to 10 percent of the maximum ',6X,' ',14X,
            &'(Sv/Bq)',12X,' ', dose',25X,' ', '
            &-----
            &' ,6X,' ',33X,' ' Weigh
            &ts Weighted Oose',7X,' ',6X,' ',33X,' ',10X,'Equivalent (Sv/Bq)
            &',2X,' ', '
            &-----

```

DOSE.FOR

10-15-1987

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&-----',/,6X,',33X,',30X,',)
DO 445 I=1,19
IF (HFIFTY(I) .EQ. 0.) THEN
  IF (NUCLID(1:2) .EQ. 'TC' .OR. (NUCLID(1:2) .EQ. 'RE')) THEN
    IF (I .EQ. 6) THEN
      IF (NDATA .EQ. 0) THEN
        WRITE(*,648) TXNAME(I), DOSE(I)
      END IF
      WRITE(65,648) TXNAME(I), DOSE(I)
448   FORMAT(6X,',',2X,A14,4X,1PE8.1,',',4X,',',30X,',')
    ELSE
      IF (NDATA .EQ. 0) THEN
        WRITE(*,650) TXNAME(I), DOSE(I)
      END IF
      WRITE(65,650) TXNAME(I), DOSE(I)
450   FORMAT(6X,',',2X,A14,4X,1PE8.1,5X,',',30X,',')
    END IF
  ELSE
    IF (NDATA .EQ. 0) THEN
      WRITE(*,650) TXNAME(I), DOSE(I)
    END IF
    WRITE(65,650) TXNAME(I), DOSE(I)
  END IF
ELSE
  IF (NUCLID(1:2) .EQ. 'TC' .OR. (NUCLID(1:2) .EQ. 'RE')) THEN
    IF (I .EQ. 6) THEN
      IF (NDATA .EQ. 0) THEN
        WRITE(*,649) TXNAME(I), DOSE(I), WDOSE(I)/HFIFTY(I), WDOSE(I)
      END IF
      WRITE(65,649) TXNAME(I), DOSE(I), WDOSE(I)/HFIFTY(I), WDOSE(I)
449   FORMAT(6X,',',2X,A14,4X,1PE8.1,',',4X,',',3X,OPF4.2,7X,1P
&E8.1,8X,',')
    ELSE
      WRITE(65,655) TXNAME(I), DOSE(I), WDOSE(I)/HFIFTY(I), WDOSE(I)
      IF (NDATA .EQ. 0) THEN
        WRITE(*,655) TXNAME(I), DOSE(I), WDOSE(I)/HFIFTY(I), WDOSE(I)
      END IF
      WRITE(65,655) TXNAME(I), DOSE(I), WDOSE(I)/HFIFTY(I), WDOSE(I)
455   FORMAT(6X,',',2X,A14,4X,1PE8.1,5X,',',3X,OPF4.2,7X,1PE8.1,8X
&,',')
    END IF
  ELSE
    WRITE(65,655) TXNAME(I), DOSE(I), WDOSE(I)/HFIFTY(I), WDOSE(I)
    IF (NDATA .EQ. 0) THEN
      WRITE(*,655) TXNAME(I), DOSE(I), WDOSE(I)/HFIFTY(I), WDOSE(I)
    END IF
  END IF
END IF
END IF
445 CONTINUE
IF (REMOD .NE. 0) THEN
  WRITE(65,660) REMOD, WTF, WREMOD
  IF (NDATA .EQ. 0) THEN
    WRITE(*,660) REMOD, WTF, WREMOD
  END IF
460  FORMAT(6X,',',2X,'Remainder',9X,1PE8.1,5X,',',3X,OPF4.2,7X,1PE8.

```

DOSE.FOR

10-15-1987

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&1,BX,'')
END IF
IF (NUCLID(1:2) .EQ. 'TC' .OR. (NUCLID(1:2) .EQ. 'RE')) THEN
  IF (NDATA .EQ. 0) THEN
    WRITE(*,462)SUM
  END IF
  WRITE(65,462)SUM
462 FORMAT(6X,'*****',14X,'-----',8X,
&'',/6X,'* + CAUTION: Stomach wall is not *',8X,'SUM = ',1PEB.1,
&8X,'',/6X,'* included as a source organ *',14X,'-----',8
&X,'',/6X,'*****',
&'*****',/6X,'',64X,'')
ELSE
  IF (NDATA .EQ. 0) THEN
    WRITE(*,461)SUM
  END IF
  WRITE(65,461)SUM
461 FORMAT(6X,'',33X,'',14X,'-----',8X,'',/6X,'',33X,'',8X,'S
&UM = ',1PEB.1,8X,'',/6X,'',33X,'',14X,'-----',8X,'',/6X,'
&'*****',
&'',/6X,'',64X,'')
END IF
IF (POST .NE. ALI) THEN
  WRITE(65,465)POST,ALI,TNAME(IRGANT)
  IF (NDATA .EQ. 0) THEN
    WRITE(*,465)POST,ALI,TNAME(IRGANT)
  END IF
465 FORMAT(6X,'',5X,'Stochastic Risk',8X,'Non-Stochastic Risk (Orga
&n)',9X,'',/6X,'',4X,1PEB.1,' Bq ',14X,1PEB.1,' Bq ',2X,'( ',A14,
&' )',4X,'')
ELSE
  WRITE(65,470)ALI
  IF (NDATA .EQ. 0) THEN
    WRITE(*,470)ALI
  END IF
470 FORMAT(6X,'',5X,'Annual Limit on Intake = ',1PEB.1,' Bq',20X
&,'')
END IF
IF (OPT .EQ. 2) THEN
  WRITE(65,475)DAC
  IF (NDATA .EQ. 0) THEN
    WRITE(*,475)DAC
  END IF
475 FORMAT(6X,'',5X,'Derived Air Concentration = ',1PEB.1,' Bq/m^3'
&,'',16X,'')
END IF
IF (NDATA .EQ. 0) THEN
  WRITE(*,477)
END IF
WRITE(65,477)
477 FORMAT(6X,'',64X,'',/,' *****
&'*****')
480 IF (NDATA .EQ. 1) GOTO 255
GOTO 125

```

DOSE.FOR

10-15-1987

```
1000 IF (DOSERR(2) .GT. 0) THEN
  WRITE (*,485)
485  FORMAT(//,' ERROR: The mode "Submerseion" can only be chosen for
    & noble gases, and elemental tritium!',//, ' SOURCE: DOSE FORTRAN',/
    &,' CORRECTIVE ACTION: Check the symbol and try again !'///// )
  DOSERR(2)=0
  IF (NDATA .NE. 0) GOTO 9009
  PAUSE ' TO RESUME PRESS <RETURN>!!'
  GOTO 125
ELSE IF (DOSERR(3) .GT. 0) THEN
  WRITE (*,490)
490  FORMAT(//,' ERROR: The entered symbol could not be found in the
    & chart of nuclides for atomic number!',//, ' SOURCE: DOSE FORTRAN',/
    &,' CORRECTIVE ACTION: Try again !'///// )
  DOSERR(3)=0
  IF (NDATA .NE. 0) GOTO 9009
  PAUSE ' TO RESUME PRESS <RETURN>!!'
  GOTO 125
END IF
9000 REWIND 5
9001 CALL CLEAR
  PRINT *, 'Do you really wish to terminate the program?'
  PRINT *
  PRINT *, ' 1 Yes'
  PRINT *, ' 2 No'
  READ(*,115,ERR=9001,END=9000)IDO
  IF (IDO .NE. 1 .AND. (IDO .NE. 2)) GOTO 9000
  IF (IDO .ED. 1) THEN
9009  STOP ' PROGRAM IS TERMINATED!!!'
  ELSE
    GOTO 125
  END IF
END
* CHARACTER*32 FUNCTION UCASE(A,M,N)
* EXAMINES STRING A, CONVERTING N CHARACTERS, STARTING WITH CHARACTER M
* TO UPPER CASE
* CHARACTER*32 A
* CHARACTER*26 LC,UC
* DATA LC/'abcdefghijklmnopqrstuvwxyz'/
* DATA UC/'ABCDEFGHIJKLMNOPQRSTUVWXYZ'/
* DO 9010 I=0,N-1
* DO 9010 J=1,26
* IF (A(M+I:M+1) .ED. LC(J:J)) THEN
* A(M+I:M+1)=UC(J:J)
* UCASE=A
* RETURN
* END IF
*9010 CONTINUE
* END
```

ENERGY, FOR

10-15-1987

```
*****
*
*      SUBROUTINE NAME: ENERGY FORTRAN
*      PURPOSE: Gives an upper and lower bound on energy of gamma
*              to help interpolate the absorbed fraction in tissue
*
*****

SUBROUTINE ENERGY(E,ELO,ENI,ILO)
  IF (E .LE. 0.010)THEN
    ILO=1
    ELO=0.010
    ENI=0.015
  ELSE IF (E .GT. 0.010 .AND. E .LE. 0.015)THEN
    ILO=1
    ELO=0.010
    ENI=0.015
  ELSE IF (E .GT. 0.015 .AND. E .LE. 0.020)THEN
    ILO=2
    ELO=0.015
    ENI=0.020
  ELSE IF (E .GT. 0.020 .AND. E .LE. 0.030)THEN
    ILO=3
    ELO=0.020
    ENI=0.030
  ELSE IF (E .GT. 0.030 .AND. E .LE. 0.050)THEN
    ILO=4
    ELO=0.030
    ENI=0.050
  ELSE IF (E .GT. 0.050 .AND. E .LE. 0.100)THEN
    ILO=5
    ELO=0.050
    ENI=0.100
  ELSE IF (E .GT. 0.100 .AND. E .LE. 0.200)THEN
    ILO=6
    ELO=0.100
    ENI=0.200
  ELSE IF (E .GT. 0.200 .AND. E .LE. 0.500)THEN
    ILO=7
    ELO=0.200
    ENI=0.500
  ELSE IF (E .GT. 0.500 .AND. E .LE. 1.000)THEN
    ILO=8
    ELO=0.500
    ENI=1.000
  ELSE IF (E .GT. 1.000 .AND. E .LE. 1.500)THEN
    ILO=9
    ELO=1.000
    ENI=1.500
  ELSE IF (E .GT. 1.500 .AND. E .LE. 2.000)THEN
    ILO=10
    ELO=1.500
    ENI=2.000
  ELSE IF (E .GT. 2.000 .AND. E .LE. 4.000)THEN
    ILO=11
```

ENERGY.FOR

10-15-1987

```
ELO=2.000  
EHI=4.000  
ELSE  
ILO=11  
ELO=2.000  
EHI=4.000  
END IF  
RETURN  
END
```

F1VALU.FOR

10-15-1987

```
*****
*
*      SUBROUTINE NAME: F1VALU FORTRAN
*
*      PURPOSE: Fractional transfer of element from GI system to
*               body fluids
*
*      Auxiliary function subprogram required: FRAC FORTRAN
*
*****
SUBROUTINE F1VALU(KZ,F1,*)
F1=0.
F1=FRAC(KZ,ITRACK)
IF (ITRACK .EQ. 1) THEN
  PAUSE ' TO RESUME PRESS <RETURN>'
  RETURN 1
END IF
RETURN
END
```


FACTOR.FOR

10-15-1987

```
*****
*
*      SUBROUTINE NAME : FACTOR  FORTRAN
*      PURPOSE: Generate matrix of specific effective
*               energy for prescribed radionuclide
*      AUXILIARY PROGRAMS REQUIRED:  a) SPEFF  FORTRAN
*                                   b) DECAY  FORTRAN
*                                   c) INTRPT  FORTRAN
*                                   d) ENERGY FORTRAN
*
*****
SUBROUTINE FACTOR (WORD,NU,SFACT,NDATA,NUCLID,PLIFE,*,*)
  DIMENSION SFACT(19,18),TMASS(19),UF(3)
  CHARACTER*8 WORD,NUCLID
  DATA UF/1.,2.131,0.5759/
  DATA TMASS/999.,19.6,37.1,8.27,1500.,150.,1040.,209.,160.,1810.,28
&4.,45.1,48200.,10500.,2830.,174.,65.4,60.3,69900./
  DO 15 I=1,19
    DO 15 J=1,17
      SFACT(I,J)=UF(NU)*SPEFF(WORD,J,I,TMASS(I),0,NUCLID,PLIFE)
    IF (WORD(1:5) .EQ. 'SORRY') THEN
      IF (NDATA .EQ. 0) THEN
        PAUSE ' TO RESUME PRESS <RETURN>!'
        RETURN 1
      ELSE
        RETURN 2
      END IF
    END IF
  15 CONTINUE
  DO 20 I=1,19
    20 SFACT(I,18)=SFACT(I,12)
    TEMP1=SFACT(5,12)
    TEMP2=SFACT(14,12)
    SFACT(5,12)=TEMP1+UF(NU)*SPEFF(WORD,12,5,TMASS(5),1,NUCLID,PLIFE)
    SFACT(5,18)=TEMP1+UF(NU)*SPEFF(WORD,12,5,TMASS(5),2,NUCLID,PLIFE)
    SFACT(14,12)=TEMP2+UF(NU)*SPEFF(WORD,12,14,TMASS(14),1,NUCLID,PLIF
&E)
    SFACT(14,18)=TEMP2+UF(NU)*SPEFF(WORD,12,14,TMASS(14),2,NUCLID,PLIF
&E)
  RETURN
END
```

FRAC.FOR

10-15-1987

```
*****
*
*      FUNCTION SUBPROGRAM NAME: FRAC FORTRAN
*      PURPOSE: Retrieve fractional transfer of isotope to the
*               body fluid compartment, F1
*      DATA FILE REQUIRED: a) BFFRAC FILE
*
*****
      FUNCTION FRAC (KZ,ITRACK)
      CHARACTER*31 C1
      OPEN(UNIT=14,FILE='BFFRAC',ACCESS='DIRECT',FORM='FORMATTED',RECL=3
&9)
      ISAVE=(5*(KZ-1))
*****
*
*      The user chooses the proper F1 according to the ingestion
*      form or inhalation class
*
*****
1 CALL CLEAR
  WRITE (*,5)
  WRITE (*,6)
5 FORMAT(///,' Enter the appropriate value of F1 from the given choi
&CES:',//)
6 FORMAT('  F1      INGESTION FORM  INHALATION CLASS',/)
  F=0.
  DO 2 I=1,5
    KEY=ISAVE+I
    READ(UNIT=14,FMT='(E8.1,A31)',ERR=10,REC=KEY)B,C1
    IF (I .EQ. 1 .AND. (B .EQ. 0))GOTO 10
    IF (B .EQ. 0)GOTO 2
    WRITE (UNIT=*,FMT='(F8.5,3X,A31,3X)',)B,C1
2 CONTINUE
4 READ(*,*,ERR=1,END=1)F
  FRAC=F
  RETURN
*****
*
*      ERROR HANDLER
*
*****
10 CALL CLEAR
  WRITE (*,15)
15 FORMAT(///,' ERROR: Value of F1 not found in the catalogue of nuclei
&des in ICRP Publication 30',/,,' SOURCE: FRAC FORTRAN',/,,' CORRECT:
&VE ACTION: Try again !!',/////))
  ITRACK=1
  RETURN
END
```

I1.FOR

10-15-1987

```
*****
*
*      FUNCTION SUBPROGRAM NAME: I1 FORTRAN
*      PURPOSE: Convert the source organ name in alphameric
*               characters to an integer from source list
*
*****

      FUNCTION I1(C2)
      CHARACTER*20 C2
      IF (C2 .EQ. ' KIDNEYS          ') THEN
         I1=6
      ELSE IF (C2 .EQ. ' LIVER          ') THEN
         I1=7
      ELSE IF (C2 .EQ. ' OVARIES        ') THEN
         I1=10
      ELSE IF (C2 .EQ. ' PANCREAS       ') THEN
         I1=11
      ELSE IF (C2 .EQ. ' MINERAL BONE   ') THEN
         I1=12
      ELSE IF (C2 .EQ. ' SPLEEN         ') THEN
         I1=14
      ELSE IF (C2 .EQ. ' TESTES         ') THEN
         I1=15
      ELSE IF (C2 .EQ. ' TNYROID        ') THEN
         I1=16
      ELSE IF (C2 .EQ. ' TOTAL BODY     ') THEN
         I1=17
      ELSE IF (C2 .EQ. ' ALL OTHER      ') THEN
         I1=18
      ELSE IF (C2 .EQ. ' BRAIN          ') THEN
         I1=19
      ELSE IF (C2 .EQ. ' ADRENALS       ') THEN
         I1=20
      ELSE IF (C2 .EQ. ' RED MARROW     ') THEN
         I1=21
      END IF
      RETURN
      END
```

ICLASS.FOR

10-15-1987

```
*****
*
*      SUBROUTINE NAME : ICLASS FORTRAN
*      PURPOSE: Find inhalation class of the given radionuclide
*      Auxiliary subroutine required: PCLASS FORTRAN
*
*****
      SUBROUTINE ICLASS(CLASS)
      CHARACTER*1 CLASS
*      CHARACTER*32 UCASE
1  CALL CLEAR
   PRINT *
   PRINT *, '          INHALATION CLASS'
   PRINT *, '          -----'
   PRINT *, ' CLASS Y---> AVID RETENTION: cleared slowly (years)'
   PRINT *, ' CLASS W---> MODERATE RETENTION: intermediate clearance (
&weeks)'
   PRINT *, ' CLASS D---> MINIMAL RETENTION: rapid clearance (days)'
   PRINT *
   PRINT *
   PRINT *, ' If you wish to examine recommendations aiding selection'
   PRINT *, ' of the pulmonary clearance classification,'
   PRINT *, ' ENTER 1; otherwise ENTER 0.'
   PRINT *
   READ (*,5,ERR=1,END=1)IWISN
5  FORMAT(I1)
   IF (IWISN.NE.1.AND.(IWISN.NE.0))GOTO 1
   IF (IWISN.EQ.1)THEN
      CALL PCLASS(IWISN)
   END IF
10 CALL CLEAR
   PRINT *
   PRINT *, ' Now ENTER the inhalation class of the given radionuclide
& (D,W, or Y)'
   PRINT *
   READ (*,15,ERR=10,END=10)CLASS
15 FORMAT(A1)
*   CLASS=UCASE(CLASS,1,1)
   IF (CLASS.NE.'D'.AND.(CLASS.NE.'W').AND.(CLASS.NE.'Y'))G
&OTO 10
   RETURN
   END
```

ICRP.FOR

10-15-1987

```

*****
*
*   SUBROUTINE NAME : ICRP FORTRAN
*   PURPOSE: Call appropriate subroutine for dose commitments
*           according to the mode of intake
*
*   DESCRIPTION OF VARIABLES
*   -----
*   INTAKE ---> 1: Ingestion, 2: inhalation, 3: submersion
*   WORD ----> Name of the given isotope, e.g., IN-113M
*   SEX ----> M or F
*   F1 ----> Fractional transfer, GI to body fluids
*   CLASS----> Pulmonary uptake classification
*   AMAD ----> Activity median aerodynamic diameter(micrometer)
*   ROB ----> Mass of 'other tissue'
*   KZ ----> Atomic number of the given nuclide
*   US ----> Matrix of transformations of nuclide, i in
*           source organ, j
*   HFIFTY----> Specific committed dose equivalent to target
*           organ or tissue
*
*****
SUBROUTINE ICRP(INTAKE,WORD,SEX,F1,CLASS,AMAD,KZ,HFIFTY,US,ROB,NDA
&TA,DER,RISK,ORGAN,*,*)
DIMENSION HFIFTY(1:24),FNP(1:20),FTB(1:20),FP(1:20),US(1:20,1:50)
CHARACTER*1 SEX,CLASS
CHARACTER*8 WORD
INTEGER ORGAN
*****
*
*   Initializing HFIFTY to be zero
*
*****
DO 5 I=1,19
HFIFTY(I)=0.
5 CONTINUE
DER=0.
RISK=0.
ORGAN=0
*****
*
*   Ingestion
*
*****
IF (INTAKE .EQ. 1)THEN
CALL INGEST(WORD,KZ,SEX,F1,HFIFTY,ROB,US,*15)
*****
*
*   Inhalation
*
*****
ELSE IF (INTAKE .EQ. 2)THEN
CALL INHALE(WORD,KZ,SEX,CLASS,F1,HFIFTY,FNP,FTB,FP,ROB,US,*15)
*****

```

ICRP.FOR

10-15-1987

```
*
*
*      Submersion
*
*****
      ELSE IF (INTAKE .EQ. 3) THEN
        CALL SUBMER(WORD,HFIFTY,DER,RISK,ORGAN,*15)
      END IF
*****
*
*      Particle Size Correction
*
*****
      IF (AMAD .NE. 1.) THEN
        IF (AMAD .LT. 0.2) THEN
          DTB=-0.163-(0.151*LOG(AMAD))
          DNP=-0.059-(0.068*LOG(AMAD))
          DP=0.289-(0.126*LOG(AMAD))
        ELSE IF (AMAD .GE. 0.2 .AND. (AMAD .LT. 10)) THEN
          DTB=0.08
          DNP=0.351+(0.219*LOG(AMAD))
          DP=0.289-(0.126*LOG(AMAD))
        ELSE
          DTB=0.229-(0.065*LOG(AMAD))
          DNP=0.621+(0.110*LOG(AMAD))
          DP=0.141-(0.040*LOG(AMAD))
        END IF
        DD 10 I=1,19
      10  HFIFTY(I)=HFIFTY(I)*((FNP(I)*DNP/0.3)+(FTB(I)*DTB/0.08)+(FP(I)*D
        &P/0.25))
      END IF
      RETURN
15  IF (NDATA .NE. 0) THEN
      RETURN 2
    ELSE
      RETURN 1
    END IF
  END
```

INGEST.FOR

10-15-1987

```

*****
*
*      SUBROUTINE NAME: INGEST FORTRAN
*
*      PURPOSE: Calculate Specific Committed Dose Equivalent (Sv/Bq)*
*               in target organs from the ingested radionuclide
*
*      AUXILIARY SUBROUTINES REQUIRED:
*
*          a) FRAC FORTRAN
*          b) DECAY1 FORTRAN
*          c) THALF FORTRAN
*          d) REPMAN FORTRAN
*          e) TFRAC FORTRAN
*          f) TRNSFM FORTRAN
*          g) SPEFF FORTRAN
*
*      DATA FILES REQUIRED:
*
*          a) EXCEPT FILE
*          b) INDEXD FILE
*
*****
*
*      DESCRIPTION OF VARIABLES:
*
*
*          WORD---->Name of the given isotope
*          KZ----->Atomic Number
*          NFIFTY-->Specific Committed Dose Equivalent (Sv/Bq)
*          ND----->Number of daughters plus one (for the parent)
*          RHALF--->Vector of half-lives of the given isotope and
*                   its daughters
*          ULIFE--->Vector of half-life units of the given isotope
*                   and its daughters
*          BRA----->Vector of branching ratios of the given isotope
*                   (BRA=1), and its daughters
*          RADID--->Vector of names of the given isotope, and its
*                   daughters
*
*****
*
*      SUBROUTINE INGEST (WORD,KZ,SEX,F1,NFIFTY,ROB,US,")
*      DIMENSION RHALF(1:50),BRA(1:50),F2(1:3),BHALF(1:3),NFIFTY(1:24),F
*      &(1:50),RCONST(1:50),AST(1:50),ASI(1:50),US(1:20,1:50),UROB(1:50),F
*      &GI(1:50)
*      CHARACTER*1 ULIFE(50),SEX
*      CHARACTER*8 RADID(50),WORD,ISOTOP,ERT,MOTS
*      REAL MROB
*      NO=0
*
*****
*
*      Subroutine for half-lives and names of the given parent
*      isotope and its daughters
*
*      DECAY MODE : A---->B---->C---->
*
*****
*
*      CALL DECAY1(WORD,RHALF,ULIFE,BRA,RADIO,NO,"12)
*      MOTS=WORD
*      DO 5 I=1,ND
*      ISOTOP=RADID(I)

```

INGEST.FOR

10-15-1987

```

      IF (I .NE. 1) THEN
        IF (ISOTOP(1:2) .EQ. 'AR' .OR. (ISOTOP(1:2) .EQ. 'KR') .OR. (IS
          &OTOP(1:2) .EQ. 'XE') .OR. (ISOTOP(1:2) .EQ. 'NE')) THEN
          NO=I-1
          GOTO 7
        END IF
      END IF
      5 CONTINUE
*****
*
*      Convert units of half-lives into days and calculate
*      the radiological constants
*
*****
      7 DO 10 I=1,NO
        IF (ULIFE(I) .EQ. 'S') THEN
          RHALF(I)=RHALF(I)/86400.
        ELSE IF (ULIFE(I) .EQ. 'M') THEN
          RHALF(I)=RHALF(I)/(60.*24.)
        ELSE IF (ULIFE(I) .EQ. 'H') THEN
          RHALF(I)=RHALF(I)/24.
        ELSE IF (ULIFE(I) .EQ. 'Y') THEN
          RHALF(I)=RHALF(I)*365.25
        END IF
        RCONST(I)=(LOG(2.))/RHALF(I)
      10 CONTINUE
*****
*
*      For alkaline earths (Ba, Ce, Ra, Sr), Tc, Re, Te-131, Te-132,*
*      Te-131m, Te-133, Te-133m, Te-134, and C, source-organ
*      transformations are not evaluated but retrieved directly
*      from data file "EXCEPT"
*
*****
      IF (MOTS(1:2) .EQ. 'BA' .OR. (MOTS(1:2) .EQ. 'CA') .OR. (MOTS(1:2)
        &.EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'SR') .OR. (MOTS(1:2) .EQ. 'C-')
        &.OR. (MOTS(1:2) .EQ. 'TC') .OR. (MOTS(1:2) .EQ. 'RE') .OR. (MOTS
          &.EQ. 'TE-131 ') .OR. (MOTS .EQ. 'TE-132 ') .OR. (MOTS .EQ. 'TE-1
            &31M ') .OR. (MOTS .EQ. 'TE-133 ') .OR. (MOTS .EQ. 'TE-133M ') .OR
              &. (MOTS .EQ. 'TE-134 ')) THEN
          GOTO 15
        END IF
*****
*
*      Initial activity, FT of the given radionuclide and its
*      daughters in transfer compartment
*
*****
      DO 14 I=1,NO
*****
*
*      FT of the parent (given) radionuclide
*
*****

```


INGEST. FOR

10-15-1987

```

IF (I .EQ. 1) THEN
  AST(1)=1./(24.*RCONST(1))
  IF (F1 .EQ. 1.) THEN
    FT(1)=24.*AST(1)*BRA(1)
  ELSE
    BFCNST=6.*F1/(1.-F1)
    ASI(1)=24./((24.*RCONST(1))*(6.+BFCNST+RCONST(1)))
    FT(1)=BFCNST*ASI(1)*BRA(1)
  END IF
ELSE
  *****
  *
  *      FT of the daughters
  *
  *****
  AST(1)=AST(I-1)*RCONST(1)/(24.*RCONST(1))
  IF (F1 .EQ. 1.) THEN
    FT(1)=24.*AST(1)*BRA(1)
  ELSE
    ASI(1)=((AST(I-1)*24.*RCONST(1)/((24.*RCONST(1))*(6.+BFCNST+R
&CONST(1))))*(ASI(I-1)*RCONST(1)/(6.+BFCNST+RCONST(1))))
    FT(1)=BFCNST*ASI(1)*BRA(1)
  END IF
  END IF
14  CONTINUE
  *****
  *
  *      Half life of clearance from transfer compartment
  *
  *****
  TSAVE=THALF(KZ)
  *****
  *
  *      TCONST---->The rate of loss of the stable element from the
  *                  body fluid compartment
  *      When transfer is instantaneous, to avoid an infinite
  *      quantity in the calculation of TCONST, it is assumed as
  *      zero
  *
  *****
  IF (TSAVE .EQ. 0.) THEN
    TCONST=0.
  ELSE
    TCONST=(LOG(2.))/TSAVE
  END IF
  *****
  *
  *      CALCULATION OF HFIFTY
  *
  *      Outer loop to calculate H50 in each target organ, KTARG
  *      where the target list is as follows:
  *
  *      TARGET ORGAN          KTARG NO.
  *      -----
  *

```

INGEST.FOR

10-15-1987

```

*      Lungs                      1      *
*      Thyroid                   2      *
*      Testes                    3      *
*      Ovaries                   4      *
*      Red marrow                5      *
*      Stomach wall              6      *
*      SI + contents             7      *
*      ULI well                  8      *
*      LLI well                  9      *
*      Liver                     10     *
*      Kidneys                   11     *
*      Bladder wall              12     *
*      Muscle                    13     *
*      Skeleton (BS cells)       14     *
*      Skin                      15     *
*      Spleen                    16     *
*      Uterus                    17     *
*      Pancreas                  18     *
*      Total body                19     *
*
*      Initializing the source organ transformations ee zero
*      for the given isotope and its daughters
*
*****
15 00 21 I=1,20
    DO 21 J=1,NO
      US(I,J)=0.
21 CONTINUE
    DO 22 I=1,NO
      UROB(I)=0.
22 CONTINUE
      HROB=0.
      ICONT=0
*****
DO 25 KTARG=1,19
*****
*
*      Skipping ovaries and testes ee target organs when the sex
*      of the subject is male and female respectively
*
*****
      IF (SEX .EQ. 'M') THEN
        IF (KTARG .EQ. 4) GOTO 25
      ELSE IF (SEX .EQ. 'F') THEN
        IF (KTARG .EQ. 3) GOTO 25
      END IF
*****
*
*      Calculating mass of each target organ
*
*****
      TMASS=REFMAN(KTARG)
      GRNSUM=0.
*****

```

INGEST.FOR

10-15-1987

```

*
*      Loop for adding contribution from all the sources in each
*      target organ
*
*      The source list is as follows:
*
*      SOURCE ORGAN                JSOURCE NO.
*      -----
*      Bladder content             1
*      Stomach content             2
*      SI content                  3
*      ULI content                 4
*      LLI content                 5
*      Kidneys                     6
*      Liver                      7
*      Lungs                      8
*      Muscle                     9
*      Ovaries                    10
*      Pancreas                   11
*      Mineral bone               12
*      Skin                      13
*      Spleen                    14
*      Testes                    15
*      Thyroid                   16
*      Total body                 17
*      All other                  18
*
*****
DQ 30 JSOURCE=17,1,-1
*****
*
*      Skipping ovaries and testas as source organs when the sex
*      of the subject is male and female respectively
*
*****
IF (SEX .EQ. 'M') THEN
  IF (JSOURCE .EQ. 10) GOTO 30
ELSE IF (SEX .EQ. 'F') THEN
  IF (JSOURCE .EQ. 15) GOTO 30
END IF
*****
*
*      For alkaline earths (Ba, Ca, Ra, Sr), Tc, Re, Ta-131, Ta-132,*
*      Ta-131m, Ta-133, Ta-133m, Ta-134, and C, source-organ
*      transformations are not evaluated but retrieved directly
*      from data file "EXCEPT"
*
*****
IF (MOTS(1:2) .EQ. 'BA' .OR. (MOTS(1:2) .EQ. 'CA') .OR. (MOTS(1:2)
& .EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'SR') .OR. (MOTS(1:2) .EQ. 'C-')
& .OR. (MOTS(1:2) .EQ. 'TC') .OR. (MOTS(1:2) .EQ. 'RE') .OR. (MOTS
& .EQ. 'TE-131 ') .OR. (MOTS .EQ. 'TE-132 ') .OR. (MOTS .EQ. 'TE-1
& 31M ') .OR. (MOTS .EQ. 'TE-133 ') .OR. (MOTS .EQ. 'TE-133M ') .OR
& . (MOTS .EQ. 'TE-134 ')) THEN

```

INGEST.FOR

10-15-1987

```

IF (KTARG .GT. 1)GOTO 45
IF (ICONT .EQ. 0)THEN
  OPEN (UNIT=50,FILE='INDEXD',FORM='FORMATTEO',ACCESS='DIRECT',R
&ECL=32)
  OPEN (UNIT=80,FILE='EXCEPT',FORM='FORMATTED',ACCESS='DIRECT',REC
&L=92)
  IF (MOTS(1:1) .EQ. 'T')THEN
    M1=33
  ELSE IF (MOTS(1:1) .EQ. 'S')THEN
    M1=26
  ELSE IF (MOTS(1:1) .EQ. 'R')THEN
    M1=14
  ELSE IF (MOTS(1:1) .EQ. 'C')THEN
    M1=9
  ELSE
    M1=1
  END IF
  DO 31 IND=M1,46
    READ(50,34,REC=IND,ERR=70)ERT,T,IREEOD,U,IREEOD
34  FORMAT(A8,F8.5,14,F8.5,14)
    IF (MOTS .EQ. ERT)THEN
      IF (T .EQ. F1)THEN
        ICONT=IREEOD
        GOTO 36
      ELSE IF (U .EQ. F1)THEN
        ICONT=IREEOD
        GOTO 36
      END IF
    END IF
    END IF
31  CONTINUE
36  CLOSE (50)
    IF (ICONT .EQ. 0)GOTO 70
  ENO IF
  READ (80,32,REC=ICONT,ERR=75)J,(US(JSORCE,I),I=1,NO)
32  FORMAT(12,1OE9.2)
    IF (J .NE. JSORCE)THEN
      DO 33 I =1,NO
        US(JSORCE,I)=0.
33  CONTINUE
      GOTO 30
    ELSE
      ICONT=ICONT+1
    END IF
    GOTO 45
  ENDIF
*****
*
*      Initializing the fraction retained in source organ from
*      the body fluid compartment, and the biological half-life
*      of the radionuclide in source organ as zero
*
*****
    DO 35 I=1,3
      F2(I)=0.

```

INGEST.FOR

10-15-1987

```

      BHALF(1)=0.
35  CONTINUE
      SMASS=0.
*****
*
*      Skipping retention fractions for source organs stomach,
*      SI, LLI, ULI
*
*****
      IF(JSORCE .EQ. 2 .OR. (JSORCE .EQ. 3) .OR. (JSORCE .EQ. 4) .OR. (J
&SORCE .EQ. 5))GOTO 40
*****
*
*      With given KZ, the following subroutine, TFRAC will give
*      retention fraction, F2 and biological half life, BHALF
*      in source organs. If several organs, I of mass (SMASS), MI
*      are associated with different retention fractions for a
*      given KZ, then for 'total body' as a source organ, the
*      mass is taken to be 70000-(sum of MI) and the retention
*      fractions to be the ones associated with source organ,
*      'all other'
*
*****
      CALL TFRAC(KZ,BHALF,JSORCE,SMASS,*12)
*****
*
*      If a source organ does not have a unique retention fraction,
*      it is skipped because often it is included in the source
*      'total body'
*
*****
      IF (F2(1) .EQ. 0. .AND. BHALF(1) .EQ. D.)GOTO 3D
*****
*
*      This subroutine TRNSFM evaluates the source-organ
*      transformations, US in organ JSORCE for the isotope and
*      its daughters
*
*****
40  IF (KTARG .GT. 1)GOTO 45
      DO 41 JEN=1,NO
          FGI(JEN)=0.
41  CONTINUE
          IPRG=0
          CALL TRNSFM(FT,F2,BHALF,RCONST,NO,BRA,US,TCONST,JSORCE,F1,IPROG,FG
&I,SMASS,UROB,MROB,KZ)
          DO 44 I=1,NO
              IF (US(JSORCE,I) .EQ. D.)GOTO 3D
44  CONTINUE
              IF (JSORCE .EQ. 17)THEN
                  ROB=MROB
              END IF
*****
*

```

INGEST.FOR

10-15-1987

```

*      Loop to calculate product of (SEE*US) from contribution
*      of all the radionuclides (parent + daughters)
*
*****
45  Q0 50 I=1,N0
    WORD=RADIO(1)
    SEE=0.
    LOOP=0.
    IF (JSORCE .EQ. 12) THEN
      IF (KTARG .EQ. 5 .OR. (KTARG .EQ. 14)) THEN
*****
*
*      Function subprogram SPEFF calculates the specific effective
*      energy, SEE deposited in target organ, KTARG due to source
*      organ, JSORCE
*
*      Loop=0 implies photon decay from radionuclides
*
*****
      SEE=SPEFF(WORD,JSORCE,KTARG,THASS,LOOP,MOTS,RNALF(1))
      IF (WORD(1:5) .EQ. 'SORRY') THEN
        PAUSE 'TO RESUME PRESS <RETURN>!'
        RETURN 1
      END IF
      GRNSUM=GRNSUM+(US(JSORCE,I)*SEE*B6400.)
      Q0 55 LOOP=1,2
*****
*
*      Radionuclides assumed to be uniformly distributed in volume
*
*****
      IF (MOTS(1:4) .EQ. 'P-33' .OR. (MOTS(1:6) .EQ. 'NB-93M') .OR. (MOT
&(1:5) .EQ. 'NB-94') .OR. (MOTS(1:5) .EQ. 'U-232') .OR. (MOTS(1:5)
& .EQ. 'U-233') .OR. (MOTS(1:5) .EQ. 'U-234') .OR. (MOTS(1:5) .EQ.
& 'U-235') .OR. (MOTS(1:5) .EQ. 'U-236') .OR. (MOTS(1:5) .EQ. 'U-238
&') .OR. (MOTS(1:2) .EQ. 'NA') .OR. (MOTS(1:2) .EQ. 'CR') .OR. (MOT
&(1:2) .EQ. 'RB') .OR. (MOTS(1:5) .EQ. 'ZN-65') .OR. (MOTS(1:6) .E
&Q. 'PB-205') .OR. (MOTS(1:6) .EQ. 'PB-210') .OR. (MOTS(1:4) .EQ. '
&BE-7') .OR. (MOTS(1:5) .EQ. 'BE-10') .OR. (MOTS(1:4) .EQ. 'V-49')
&.OR. (MOTS(1:6) .EQ. 'PD-103') .OR. (MOTS(1:6) .EQ. 'PD-107') .OR.
& (MOTS(1:6) .EQ. 'SN-113') .OR. (MOTS(1:6) .EQ. 'SN-123') .OR. (MO
&(1:7) .EQ. 'SN-119M') .OR. (MOTS(1:6) .EQ. 'SN-126') .OR. (MOTS(
&(1:6) .EQ. 'TA-182') .OR. (MOTS(1:5) .EQ. 'W-181') .OR. (MOTS(1:5)
&.EQ. 'W-185') .OR. (MOTS(1:5) .EQ. 'W-188')) THEN
        UTRAB=0.2*US(JSORCE,I)*B6400.
        UCORT=0.8*US(JSORCE,I)*B6400.
      ELSE IF (MOTS(1:2) .EQ. 'BA' .OR. (MOTS(1:2) .EQ. 'CA') .OR. (MOT
&(1:2) .EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'SR')) THEN
        IF (RNALF(1) .GT. 15) THEN
          UTRAB=0.2*US(JSORCE,I)*B6400.
          UCORT=0.8*US(JSORCE,I)*B6400.
        ELSE
          UTRAB=0.5*US(JSORCE,I)*B6400.
          UCORT=0.5*US(JSORCE,I)*B6400.

```

INGEST.FOR

10-15-1987

```
      END IF
    ELSE
*****
*
*      Radionuclides assumed to be on bone surfaces
*
*****
      UTRAB=0.5*US(JSORCE,I)*86400.
      UCORT=0.5*US(JSORCE,I)*86400.
    END IF
*****
*
*      Loop=1 implies charged particle dose in trabecular bone
*
*****
52  IF(LOOP .EQ. 1)THEN
      SEE=SPEFF(WORD,JSORCE,KTARG,TMASS,LOOP,MOTS,RHALF(1))
      IF (WORD(1:5) .EQ. 'SORRY')THEN
        PAUSE 'TO RESUME PRESS <RETURN>!'
        RETURN 1
      END IF
      GRNSUM=GRNSUM+(UTRAB*SEE)
    ELSE
*****
*
*      Loop=2 implies charged particle dose in cortical bone
*
*****
      SEE=SPEFF(WORD,JSORCE,KTARG,TMASS,LOOP,MOTS,RHALF(1))
      IF (WORD(1:5) .EQ. 'SORRY')THEN
        PAUSE 'TO RESUME PRESS <RETURN>!'
        RETURN 1
      END IF
      GRNSUM=GRNSUM+(UCORT*SEE)
    END IF
55  CONTINUE
    ELSE
      GOTD 60
    END IF
    GOTD 50
  END IF
60  SEE=SPEFF(WORD,JSORCE,KTARG,TMASS,LOOP,MOTS,RHALF(1))
    IF (WORD(1:5) .EQ. 'SORRY')THEN
      PAUSE 'TO RESUME PRESS <RETURN>!'
      RETURN 1
    END IF
    GRNSUM=GRNSUM+(US(JSORCE,I)*SEE*86400.)
50  CONTINUE
30  CONTINUE
    HFIFTY(KTARG)=(1.6E-10)*GRNSUM
25  CONTINUE
    RETURN
12  RETURN 1
70  WRITE(*,71)
```

INGEST.FOR

10-15-1987

```
71 FORMAT(///, ' ERROR: Nuclide not found in catalogue of ICRP Publicat
&ion 30',/, ' SOURCE: INGEST FORTRAN',/, ' CORRECTIVE ACTION: Try and
&ther nuclide',///// )
      GOTO 12
75 WRITE(*,76)
76 FORMAT(///, ' ERROR: Unable to read US values from file "EXCEPT"',/
&,' SOURCE: INGEST FORTRAN',/, ' CORRECTIVE ACTION: Check the identi
&fication and try again!')
      END
```


INHALE.FOR

10-15-1987

```

*****
*
* SUBROUTINE NAME : INHALE FORTRAN
* PURPOSE: Generate e vector of committed dose equivalent
*          (Sv/Bq) in target organ due to inhaled nuclide
* AUXILIARY PROGRAMS REQUIRED: e) PCLASS FORTRAN
*                             b) DECAY1 FORTRAN
*                             c) FRAC FORTRAN
*                             d) RESPIR FORTRAN
*                             e) THALF FORTRAN
*                             f) REFMAN FORTRAN
*                             g) TFRAC FORTRAN
*                             h) TRNSFM FORTRAN
*                             i) SPEFF FORTRAN
*
* DATA FILES REQUIRED:      a) EXCEPT FILE
*                          b) INDEX1 FILE
*
*****
*
* DESCRIPTION OF VARIABLES
* -----
*
* WORD----> Name of the given isotope
* KZ-----> Atomic number
* NFIFTY--> Specific committed dose
* NO-----> Number of daughters + one for the given isotope
* RHALF--> Vector of half-lives of the given isotope and its
*          daughters
* ULIFE--> Vector of half-life units of the given isotope
*          and its daughters
* BRA-----> Vector of branching ratios of the given isotope
*          (BRA(1)=1) and its daughters
* RADIO--> Vector of names of the given isotope and its
*          daughters
*
*****
* SUBROUTINE INHALE(WORD,KZ,SEX,CLASS,F1,NFIFTY,FNP,FTG,FP,ROB,US,*)
* DIMENSION RHALF(1:50),BRA(1:50),F2(1:3),BNALF(1:3),NFIFTY(1:24),FT
* &(1:50),RCONST(1:50),AST(1:50),ASI(1:50),US(1:20,1:50),UROB(1:50),F
* &BF(1:50),FBFDR(1:50),AA(1:50),AB(1:50),AC(1:50),AD(1:50),AE(1:50)
* &,AF(1:50),AG(1:50),AK(1:50),AI(1:50),AD1(1:50),FGI(1:50),AJ(1:50),
* &FNP(1:20),FTB(1:20),FP(1:20),TNP(1:20,1:50),TTB(1:20,1:50),TP(1:20
* &,1:50)
* CHARACTER*1 ULIFE(50),SEX,CLASS,TYPE, SORT
* CHARACTER*8 RADIO(50),WORD,ISOTOP,ERT,MOTS
* REAL MROB
*
*****
*
* Subroutine for half-lives and names of the given isotope
* and its daughters
* DECAY MODE : A----> B----> C---->
*
*****
NO=0

```

INHALE.FOR

10-15-1987

```

CALL DECAY1(WORD,RNALF,ULIFE,BRA,RADID,NO,*12)
MDTS=WORD
DO 5 I=1,ND
  ISOTOP=RADIO(I)
  IF (I .EQ. 1) THEN
    IF (ISOTOP(1:2) .EQ. 'AR' .OR. (ISOTOP(1:2) .EQ. 'KR') .OR. (IS
&OTDP(1:2) .EQ. 'XE') .OR. (ISOTOP(1:2) .EQ. 'NE')) THEN
      NO=I-1
      GOTD 7
    END IF
  END IF
5 CONTINUE
*****
*
*   Convert units of half-lives into days and calculate the
*   radiological constants
*
*****
7 DO 10 I=1,NO
  IF (ULIFE(I) .EQ. 'S') THEN
    RNALF(I)=RNALF(I)/86400.
  ELSE IF (ULIFE(I) .EQ. 'H') THEN
    RNALF(I)=RNALF(I)/(60.*24.)
  ELSE IF (ULIFE(I) .EQ. 'M') THEN
    RNALF(I)=RNALF(I)/24.
  ELSE IF (ULIFE(I) .EQ. 'Y') THEN
    RNALF(I)=RNALF(I)*365.25
  END IF
  RCONST(I)=(LOG(2.))/FLOW(RNALF(I))
10 CONTINUE
*****
*
*   Fraction of inhaled stable element transferred to the
*   body fluids via the GI tract, FBF
*
*****
DD 15 I=1,ND
*****
*
*   FBF of the parent (given) radionuclide
*
*****
IF (I .EQ. 1) THEN
  AST(1)=1./FLOW(24.*RCONST(1))
  IF (F1 .EQ. 1.) THEN
    FBF(1)=FLOW(24.*AST(1)*BRA(1))
  ELSE
    BFCNST=.6.*F1/(1.-F1)
    ASI(1)=24./FLOW((24.*RCONST(1))*(6.+BFCNST+RCONST(1)))
    FBF(1)=FLOW(BFCNST*ASI(1)*BRA(1))
  END IF
ELSE
*****
*

```

INHALE.FOR

10-15-1987

```

*          FBF of the daughters
*
*****
AST(I)=AST(I-1)*RCONST(1)/FLOW(24.*RCONST(1))
IF (F1.EQ.1.)THEN
    FBF(I)=FLOW(24.*AST(I)*BRA(I))
ELSE
    ASI(I)=(AST(I-1)*24.*RCONST(1)/FLOW((24.*RCONST(1))*6.*BFCN
&ST+RCONST(1)))+(ASI(I-1)*RCONST(1)/FLOW(6.*BFCNST+RCONST(1)))
    FBF(I)=FLOW(BFCNST*ASI(I)*BRA(I))
END IF
END IF
15    CONTINUE
*****
*
*          Fractions of inhaled material deposited in three respiratory
*          regions, the nasal passage (N-P), the trachea and bronchial
*          tree (T-B), and the pulmonary region (P), the balance being
*          the fraction exhaled. It is assumed that the activity median
*          aerodynamic diameter, AMAD is 1 micrometer
*
*****
DNP=0.30
DTB=0.08
DP=0.25
*****
*
*          Subroutine for fraction and clearance rates for transfer of
*          the material between compartments
*          Initializing all clearance rates and fractions to zero
*
*****
DATA FA,FB,FC,FD,FE,FF,FG,FH,FI,FJ /10*0./
DATA CLA,CLB,CLC,CLD,CLE,CLF,CLG,CLH,CLI,CLJ /10*0./
CALL RESPIR (CLASS,FA,FB,FC,FD,FE,FF,FG,FH,FI,FJ,CLA,CLB,CLC,CLD,C
&LE,CLF,CLG,CLH,CLI,CLJ)
*****
*
*          Transformations in various compartments of the lung
*
*****
DO 25 I=1,NO
*****
*
*          Transformations of the parent radionuclide
*
*****
IF (I.EQ.1)THEN
    AA(1)=DNP*FA/FLOW(CLA+RCONST(1))
    AB(1)=DNP*FB/FLOW(CLB+RCONST(1))
    AC(1)=DTB*FC/FLOW(CLC+RCONST(1))
    AD(1)=DTB*FD/FLOW(CLD+RCONST(1))
    AE(1)=DP*FE/FLOW(CLE+RCONST(1))
    AH(1)=DP*FH/FLOW(CLH+RCONST(1))

```

INHALE.FOR

10-15-1987

```

AI(1)=AH(1)*CLH*FI/FLOW(CLI+RCONST(1))
AJ(1)=0.
IF (CLASS .EQ. 'D') THEN
  AD1(1)=0.
  AF(1)=0.
  AG(1)=0.
ELSE
  AF(1)=DP*FF/FLOW(CLF+RCONST(1))
  AG(1)=DP*FG/FLOW(CLG+RCONST(1))
  AD1(1)=(AF(1)*CLF)+(AG(1)*CLG)/FLOW(CLD+RCONST(1))
END IF
IF (CLASS .EQ. 'Y') THEN
  AJ(1)=DP*FH*CLH*FJ*(1.-UXP(-365.25*50.*RCONST(1)))/FLOW(RCON
&ST(1)*(CLH+RCONST(1)))
END IF
ELSE
*****
*
*      Transformations of the daughters
*
*****
AA(1)=AA(I-1)*RCONST(1)/FLOW(CLA+RCONST(1))
AB(1)=AB(I-1)*RCONST(1)/FLOW(CLB+RCONST(1))
AC(1)=AC(I-1)*RCONST(1)/FLOW(CLC+RCONST(1))
AD(1)=AD(I-1)*RCONST(1)/FLOW(CLD+RCONST(1))
AE(1)=AE(I-1)*RCONST(1)/FLOW(CLE+RCONST(1))
AH(1)=AH(I-1)*RCONST(1)/FLOW(CLH+RCONST(1))
AI(1)=(AH(1)*CLH*FI/FLOW(CLI+RCONST(1)))+(AI(I-1)*RCONST(1)/FLO
&W(CLI+RCONST(1)))
AJ(1)=0.
IF (CLASS .EQ. 'D') THEN
  AD1(1)=0.
  AF(1)=0.
  AG(1)=0.
ELSE
  AF(1)=AF(I-1)*RCONST(1)/FLOW(CLF+RCONST(1))
  AG(1)=AG(I-1)*RCONST(1)/FLOW(CLG+RCONST(1))
  AD1(1)=(AF(1)*CLF)+(AG(1)*CLG)/FLOW(CLD+RCONST(1))
END IF
IF (CLASS .EQ. 'Y') THEN
  AJ(1)=(AJ(I-1)+(AH(I-1)*CLH*FI/FLOW(CLH+RCONST(1))))*(1.-UXP
&(-365.25*50.*RCONST(1)))
END IF
END IF
25 CONTINUE
*****
*
*      Fraction of the inhaled radionuclide transferred directly to
*      the body fluid compartment, FBFDIR
*
*****
DO 27 I=1,NO
  FBFDIR(I)=BRA(I)*FLOW((CLA*AA(I))+(CLC*AC(I))+(CLE*AE(I))+(CLI*AI(
&I)))

```

INHALE.FOR

10-15-1987

27 CONTINUE

```
*****
*
* Fraction of the inhaled radionuclide transferred to the
* gastro-intestinal tract, FGI
*
*****
```

```
DO 30 I=1,N0
  FGI(I)=BRA(I)*FLOW((CLB*AB(I))+ (CLD*AD(I))+ (CLO*AD1(I)))
30 CONTINUE
```

```
*****
*
* Total initial activity, FT of the given radionuclide and
* its daughters in the TRANSFER compartment
*
*****
```

```
DO 35 I=1,N0
  FT(I)=FBFDIR(I)+(FGI(I)*FBF(I))
35 CONTINUE
```

```
*****
*
* For alkaline earths (Be, Ca, Ra, Sr), Tc, Ra, Te-131, Te-132,
* Te-131m, Te-133, Te-133m, Te-134, and C, source-organ
* transformations are not evaluated but retrieved directly
* from data file "EXCEPT"
*
*****
```

```
IF (MOTS(1:2) .EQ. 'BA' .OR. (MOTS(1:2) .EQ. 'CA') .OR. (MOTS(1:2)
& .EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'SR') .OR. (MOTS(1:2) .EQ. 'C-')
& .OR. (MOTS(1:2) .EQ. 'TC') .OR. (MOTS(1:2) .EQ. 'RE') .OR. (MOTS(
& 1:6) .EQ. 'TE-131') .OR. (MOTS(1:6) .EQ. 'TE-132') .OR. (MOTS(1:7)
& .EQ. 'TE-131M') .OR. (MOTS(1:6) .EQ. 'TE-133') .OR. (MOTS(1:7) .E
& .EQ. 'TE-133M') .OR. (MOTS(1:6) .EQ. 'TE-134')) THEN
```

GOTO 37

END IF

```
*****
*
* Half-life of clearance from the TRANSFER compartment
*
*****
```

TSAVE=HALF(K2)

```
*****
*
* TCONST---- The rate of loss of the stable element from the
* body fluid compartment
* When transfer is instantaneous, to avoid an infinite quantity
* in the calculation of TCONST, it is assumed as zero
*
*****
```

IF (TSAVE .EQ. 0.) THEN

TCONST=0.

ELSE

TCONST=(LOG(2.))/TSAVE

END IF

INHALE.FOR

10-15-1987

```
*****
*
*          CALCULATION OF HFIFTY
*-----*
*          TARGET LIST
*-----*
*          TARGET ORGAN          KTARG NO.
*-----*
*          Lungs                  1
*          Thyroid                2
*          Testes                 3
*          Ovaries                4
*          Red marrow             5
*          Stomach wall           6
*          SI + contents          7
*          ULI wall               8
*          LLI wall               9
*          Liver                  10
*          Kidneys                11
*          Bladder wall           12
*          Muscles                13
*          Skeleton (BS cells)    14
*          Skin                   15
*          Spleen                 16
*          Uterus                 17
*          Pancreas               18
*          Total body             19
*
*-----*
*
*          Initializing the source-organ transformations as zero for
*          the given isotope and its daughters
*
*****
37 00 40 I=1,17
    DO 40 J=1,N0
    US(I,J)=0.
40 CONTINUE
    00 42 I=1,N0
    UROB(I)=0.
42 CONTINUE
    MROB=0.
    ICONT=0
*****
*
*          Outer loop to calculate N-50 in each target organ, KTARG
*-----*
*          00 50 KTARG=1,19
*****
*
*          Skipping ovaries and testes as target organs when the sex
*          of the subject is male and female respectively
*
```

INHALE.FOR

10-15-1987

```

*****
      IF (SEX .EQ. 'M') THEN
        IF (KTARG .EQ. 4) GOTO 50
      ELSE IF (SEX .EQ. 'F') THEN
        IF (KTARG .EQ. 3) GOTO 50
      END IF
*****
*
*       Calculating mass of each target organ
*
*****
      TMASS=REFMAN(KTARG)
      GRNSUM=0.
      FNP(KTARG)=0.
      FTB(KTARG)=0.
      FP(KTARG)=0.
      DO 53 I=1,17
      DO 53 J=1,NO
        TMP(I,J)=0.
        TTB(I,J)=0.
        TP(I,J)=0.
      53 CONTINUE
*****
*
*       Loop for adding contribution from all the sources in each
*       target organ
*
*-----*
*       THE SOURCE LIST IS AS FOLLOWS:
*-----*
*
*       SOURCE ORGAN                JSORCE NO.
*       -----
*       Bladder content              1
*       Stomach content              2
*       SI content                   3
*       ULI content                  4
*       LLI content                  5
*       Kidneys                      6
*       Liver                        7
*       Lunga                        8
*       Muscle                       9
*       Ovaries                      10
*       Pancreas                    11
*       Mineral bone                12
*       Skin                        13
*       Spleen                      14
*       Testes                      15
*       Thyroid                     16
*       Total body                   17
*       All other                    18
*
*****
      DO 55 JSORCE=17,1,-1
*****

```

INHALE.FOR

10-15-1987

```

*
*      Skipping ovaries and testes as source organs when the sex of
*      the subject is male and female respectively
*
*****
      IF (SEX .EQ. 'M') THEN
        IF (JSORCE .EQ. 10) GOTD 55
      ELSE IF (SEX .EQ. 'F') THEN
        IF (JSORCE .EQ. 15) GOTD 55
      END IF
*****
*
*      For alkali earths (Be, Ca, Re, Sr), Tc, Re, Ta-131, Te-132,
*      Te-131m, Ta-133, Te-133m, Te-134, and C, source organ
*      transformations are not evaluated but retrieved directly
*      from data file "EXCEPT"
*
*****
      IF (MOTS(1:2) .EQ. 'BA' .OR. (MOTS(1:2) .EQ. 'CA') .OR. (MOTS(1:2)
& .EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'SR') .OR. (MOTS(1:2) .EQ. 'C-')
& .OR. (MOTS(1:2) .EQ. 'TC') .OR. (MOTS(1:2) .EQ. 'RE') .OR. (MOTS(
& 1:6) .EQ. 'TE-131') .OR. (MOTS(1:6) .EQ. 'TE-132') .OR. (MOTS(1:7)
& .EQ. 'TE-131m') .OR. (MOTS(1:6) .EQ. 'TE-133') .OR. (MOTS(1:7) .E
& .EQ. 'TE-133m') .OR. (MOTS(1:6) .EQ. 'TE-134')) THEN
        IF (KTARG .GT. 1) GOTD 70
        IF (ICONT .EQ. 0) THEN
          OPEN(UNIT=50, FILE='INDEX1', FORM='FORMATTED', ACCESS='DIRECT',
& RECL=34)
          OPEN(UNIT=80, FILE='EXCEPT', FORM='FORMATTED', ACCESS='DIRECT',
& RECL=92)
          IF (MOTS(1:1) .EQ. 'T') THEN
            M1=33
          ELSE IF (MOTS(1:1) .EQ. 'S') THEN
            M1=26
          ELSE IF (MOTS(1:1) .EQ. 'R') THEN
            M1=14
          ELSE IF (MOTS(1:1) .EQ. 'C') THEN
            M1=9
          ELSE
            M1=1
          END IF
          DO 56 IND=M1,46
            READ(50,57,REC=IND,ERR=100) ERT,T,TYPE,IREFCO,U,SORT,IREFCO
            FORMAT(A8,F8.5,A1,I4,F8.5,A1,I4)
57      IF (MOTS .EQ. ERT) THEN
              IF (T .EQ. F1 .AND. (TYPE .EQ. CLASS)) THEN
                ICONT=IREFCO
                GOTD 58
              ELSE IF (U .EQ. F1 .AND. (SORT .EQ. CLASS)) THEN
                ICONT=IREFCO
                GOTD 58
              END IF
            END IF
56      CONTINUE

```


INHALE.FOR

10-15-1987

```

58      CLOSE (50)
         IF (ICOMT.EQ. 0)GOTO 100
      END IF
         IF (ICOMT.EQ. 696)GOTO 70
      READ(80,59,REC=ICOMT,ERR=105)J,(US(JSORCE,I),I=1,NO)
59      FORMAT(12,10E9.2)
         IF (J.NE. JSORCE)THEN
            DO 61 I=1,NO
               US(JSORCE,I)=0.
61      CONTINUE
            GOTO 55
         ELSE
            ICOMT=ICOMT+1
      END IF
      GOTO 70
      END IF

*****
*
*
*      Initializing the fraction retained in source organ from the
*      body fluid compartment, and the biological half-life of the
*      radionuclide in source organ ee zero
*
*****
      DO 60 I=1,3
         F2(I)=0.
         BNALF(I)=0.
60 CONTINUE
         SMASS=0.

*****
*
*      Skipping retention fractions for source organs stomach,
*      SI, LLI, ULI, and lung
*
*****
      IF(JSORCE.EQ. 2.OR. (JSORCE.EQ. 3).OR. (JSORCE.EQ. 4).OR. (J
&SORCE.EQ. 5).OR. (JSORCE.EQ. 8))GOTO 65
*****
*
*      With given KZ, the following subroutine, TFRAC will give
*      retention fraction, F2 and biological half-life, BNALF in
*      source organs. If several organs, i of mass (SMASS) Mi are
*      associated with different retention fractions for a given KZ,
*      then for 'TOTAL BODY' as source organ, the source mass is
*      taken to be 70000-(SUM Mi) and the retention fractions to be
*      the ones associated with source organ 'ALL OTHER'
*
*****
      CALL TFRAC(KZ,F2,BNALF,JSORCE,SMASS,*12)
*****
*
*      If a source organ does not have a unique retention fraction,
*      it is skipped because often it is included in the source

```

INHALE.FOR

10-15-1987

```

*      'TOTAL BODY'
*
*****
      IF (F2(1) .EQ. 0. .AND. BHALF(1) .EQ. 0.)GOTO 55
      65 IF (KTARG .GT. 1)GOTO 70
*****
*
*      US for Respiratory system
*
*****
      IF (JSORCE .EQ. 8)THEN
      00 67 I=1,NO
      US(JSORCE,I)=BRA(1)*FLOW(AC(I)+AD(I)+AD1(I)+AE(I)+AF(I)+AG(I)+AH(I)
      &)+AI(I)+AJ(I))
      67 CONTINUE
      GOTO 70
      END IF
*****
*
*      The subroutine TRNSFM evaluates the source-organ trans-
*      formations, US in organ JSORCE for the isotope and its
*      daughters
*
*****
      IPRG=1
      CALL TRNSFM(FT,F2,BHALF,RCONST,NO,BRA,US,TCONST,JSORCE,F1,IPRG,FG
      &I,SMASH,UROB,MROB,KZ)
      00 69 I=1,NO
      IF (US(JSORCE,I) .EQ. 0.)GOTO 55
      69 CONTINUE
      IF (JSORCE .EQ. 17)THEN
      ROB=MROB
      END IF
*****
*
*      Loop to calculate product of (SEE*US) from contribution of
*      all radionuclides
*
*****
      70 00 75 I=1,NO
      WORO=RADIO(I)
      SEE=0.
      LOOP=0.
      IF (JSORCE .EQ. 12)THEN
      IF (KTARG .EQ. 5 .OR. (KTARG .EQ. 14))THEN
*****
*
*      Function subprogram SPEFF calculates the specific effective
*      energy deposited in target organ, KTARG due to source organ,
*      JSORCE
*      Loop=0 implies photon decay from radionuclides
*
*****
      SEE=SPEFF(WORO,JSORCE,KTARG,TMASS,LOOP,MOTS,RHALF(1))

```

INHALE.FOR

10-15-1987

```

IF (WORD(1:5) .EQ. 'SORRY') THEN
  PAUSE ' TO RESUME PRESS <RETURN>!!'
  RETURN 1
END IF
GRNSUM=GRNSUM+(US(JSORCE,1)*SEE*86400.)
DEPOT=US(JSORCE,1)*SEE*86400./FLOW(FT(1))
TMP(JSORCE,1)=FLOW((CLA*AA(1))+CLB*AB(1)*FBF(1))*DEPOT
TTB(JSORCE,1)=FLOW((CLC*AC(1))+CLD*AD(1)*FBF(1))*DEPOT
TP(JSORCE,1)=FLOW((CLE*AE(1))+CLF*AI(1))+CLD*AD(1)*FBF(1)
&))*DEPOT
DO 80 LOOP=1,2
*****
*
*   Radionuclides assumed to be uniformly distributed in volume
*
*****
  IF (MOTS(1:4) .EQ. 'P-33' .OR. (MOTS(1:6) .EQ. 'NB-93M' .OR. (MOT
&S(1:5) .EQ. 'NB-94' .OR. (MOTS(1:5) .EQ. 'U-232' .OR. (MOTS(1:5)
& .EQ. 'U-233' .OR. (MOTS(1:5) .EQ. 'U-234' .OR. (MOTS(1:5) .EQ.
&'U-235' .OR. (MOTS(1:5) .EQ. 'U-236' .OR. (MOTS(1:5) .EQ. 'U-238
&') .OR. (MOTS(1:2) .EQ. 'NA' .OR. (MOTS(1:2) .EQ. 'CR' .OR. (WOR
&D(1:2) .EQ. 'RB' .OR. (MOTS(1:5) .EQ. 'ZN-65' .OR. (MOTS(1:6) .E
&Q. 'PB-205' .OR. (MOTS(1:6) .EQ. 'PB-210' .OR. (MOTS(1:4) .EQ. '
&BE-7' .OR. (MOTS(1:5) .EQ. 'BE-10' .OR. (MOTS(1:6) .EQ. 'V-49'
&.OR. (MOTS(1:6) .EQ. 'PO-103' .OR. (MOTS(1:6) .EQ. 'PO-107' .OR.
& (MOTS(1:6) .EQ. 'SN-113' .OR. (MOTS(1:6) .EQ. 'SN-123' .OR. (MO
&TS(1:7) .EQ. 'SN-119M' .OR. (MOTS(1:6) .EQ. 'SN-126' .OR. (MOTS(
&1:6) .EQ. 'TA-182' .OR. (MOTS(1:5) .EQ. 'U-181' .OR. (MOTS(1:5)
&.EQ. 'U-185' .OR. (MOTS(1:5) .EQ. 'U-188')) THEN
    UTRAB=0.2*US(JSORCE,1)*86400.
    UCORT=0.8*US(JSORCE,1)*86400.
  ELSE IF (MOTS(1:2) .EQ. 'BA' .OR. (MOTS(1:2) .EQ. 'CA' .OR. (MOTS
&(1:2) .EQ. 'RA' .OR. (MOTS(1:2) .EQ. 'SR')) THEN
    IF (RHALF(1) .GT. 15) THEN
      UTRAB=0.2*US(JSORCE,1)*86400.
      UCORT=0.8*US(JSORCE,1)*86400.
    ELSE
      UTRAB=0.5*US(JSORCE,1)*86400.
      UCORT=0.5*US(JSORCE,1)*86400.
    END IF
  ELSE
*****
*
*   Radionuclides assumed to be on bone surfaces
*
*****
    UTRAB=0.5*US(JSORCE,1)*86400.
    UCORT=0.5*US(JSORCE,1)*86400.
  END IF
*****
*
*   Loop=1 implies charged particle dose in trabecular bone
*
*****

```

INHALE.FOR

10-15-1987

```

82      IF (LOOP .EQ. 1) THEN
          SEE=SPEFF(WORD,JSORCE,KTARG,TMASS,LOOP,MOTS,RHALF(1))
          IF (WORD(1:5) .EQ. 'SORRY') THEN
              PAUSE ' TO RESUME PRESS <RETURN>!!'
              RETURN 1
          END IF
          GRNSUM=GRNSUM+(UTRAB*SEE)
          DEPOT=UTRAB*SEE/FLOW(FT(1))
          TNP(JSORCE,1)=FLOW(((CLA*AA(1)))+(CLB*AB(1))*FBF(1)))*DEPOT
          &)+TNP(JSORCE,1)
          TTB(JSORCE,1)=FLOW(((CLC*AC(1)))+(CLD*AD(1))*FBF(1)))*DEPOT
          &)+TTB(JSORCE,1)
          TP(JSORCE,1)=FLOW(((CLE*AE(1)))+(CLI*AI(1)))+(CLD*AD(1))*FB
          &F(1)))*DEPOT)+TP(JSORCE,1)
          ELSE
*****
*
*      Loop=2 implies charged particle dose in cortical bone
*
*****
          SEE=SPEFF(WORD,JSORCE,KTARG,TMASS,LOOP,MOTS,RHALF(1))
          IF (WORD(1:5) .EQ. 'SORRY') THEN
              PAUSE ' TO RESUME PRESS <RETURN>!!'
              RETURN 1
          END IF
          GRNSUM=GRNSUM+(UCORT*SEE)
          DEPOT=UCORT*SEE/FLOW(FT(1))
          TNP(JSORCE,1)=FLOW(((CLA*AA(1)))+(CLB*AB(1))*FBF(1)))*DEPOT
          &)+TNP(JSORCE,1)
          TTB(JSORCE,1)=FLOW(((CLC*AC(1)))+(CLD*AD(1))*FBF(1)))*DEPOT
          &)+TTB(JSORCE,1)
          TP(JSORCE,1)=FLOW(((CLE*AE(1)))+(CLI*AI(1)))+(CLD*AD(1))*FB
          &F(1)))*DEPOT)+TP(JSORCE,1)
          END IF
80      CONTINUE
      ELSE
          GOTD 85
      END IF
      GOTD 75
      END IF
85      SEE=SPEFF(WORD,JSORCE,KTARG,TMASS,LOOP,MOTS,RHALF(1))
      IF (WORD(1:5) .EQ. 'SORRY') THEN
          PAUSE ' TO RESUME PRESS <RETURN>!!'
          RETURN 1
      END IF
      GRNSUM=GRNSUM+(US(JSORCE,1)*SEE*86400.)
*****
*
*      Fraction of committed dose equivalent in the target tissue
*      resulting from deposition in the N-P, T-B, and P regions
*
*****
      DEPOT=US(JSORCE,1)*SEE*86400.
      IF (JSORCE .EQ. 8) THEN

```

INHALE.FOR

10-15-1987

```

      TNP(JSORCE,I)=0.
      TTb(JSORCE,I)=FLOW(AC(I)+AD(I))*BRA(I)*SEE*86400.
      TP(JSORCE,I)=FLOW(AD1(I)+AE(I)+AF(I)+AG(I)+AH(I))*BRA(I)*SEE*86
      8400.
      ELSE IF(JSORCE.EQ.2.OR.(JSORCE.EQ.3).OR.(JSORCE.EQ.4)
      &.OR.(JSORCE.EQ.5))THEN
          TNP(JSORCE,I)=FLOW(CLB*AB(I))*DEPOT/FGI(I))
          TTb(JSORCE,I)=FLOW(CLD*AD(I))*DEPOT/FGI(I))
          TP(JSORCE,I)=FLOW(CLO*AD1(I))*DEPOT/FGI(I))
      ELSE
          TNP(JSORCE,I)=FLOW(((CLA*AA(I))+CLB*AB(I))*FBI(I))*DEPOT/FT
          &(I))
          TTb(JSORCE,I)=FLOW(((CLC*AC(I))+CLD*AD(I))*FBI(I))*DEPOT/FT
          &(I))
          TP(JSORCE,I)=FLOW((((CLE*AE(I))+CLI*AI(I))+CLD*AD1(I))*FBI(I)
          &)))*DEPOT/FT(I))
      END IF
75 CONTINUE
      DO 95 I=1,N0
          FNP(KTARG)=FNP(KTARG)+TNP(JSORCE,I)
          FTb(KTARG)=FTb(KTARG)+TTb(JSORCE,I)
          FP(KTARG)=FP(KTARG)+TP(JSORCE,I)
95 CONTINUE
55 CONTINUE
      GROUP=FNP(KTARG)+FTb(KTARG)+FP(KTARG)
      FNP(KTARG)=FNP(KTARG)/FLOW(GROUP)
      FTb(KTARG)=FTb(KTARG)/FLOW(GROUP)
      FP(KTARG)=FP(KTARG)/FLOW(GROUP)
*****
*
*      Specific committed dose , H-50 in each target organ
*
*****
      HFIFTY(KTARG)=(1.6E-10)*GRNSUM
50 CONTINUE
      RETURN
12 RETURN 1
100 WRITE(*,101)
101 FORMAT(//,' ERROR: Nuclide not found in catalogue of ICRP Publicat
      &ion 30',/, ' SOURCE: INHALE.FORTRAN',/, ' CORRECTIVE ACTION: Try ano
      &ther nuclide',////)
      GOTO 12
105 WRITE(*,106)
106 FORMAT(//,' ERROR: Unable to read US values from file "EXCEPT",/
      &, ' SOURCE: INHALE.FORTRAN',/, ' CORRECTIVE ACTION: Check the identi
      &fication and try again!')
      ENO

```

INTRPT.FOR

10-15-1987

```
*****
*
*   FUNCTION SUBPROGRAM NAME : INTRPT FORTRAN
*   PURPOSE: Interpolates the values of absorbed fraction
*   AUXILIARY PROGRAM REQUIRED:  e) ENERGY FORTRAN
*   DATA FILE REQUIRED:       a) ABSFRAC FILE
*
*****
      FUNCTION INTRPT (E,JSORCE,KTARG,ICHECK)
      REAL INTRPT
      OPEN (UNIT=13,FILE='ABSFRAC',ACCESS='DIRECT',FORM='FORMATTED',RECL
&=9)
      ILO=0
      ELO=0.
      ENI=0.
      CALL ENERGY(E,ELO,ENI,ILO)
      INI=ILO+1
      KEY=ILO+((JSORCE-1)*12)+((KTARG-1)*12*20)
      READ (UNIT=13,FMT='(E9.3)',REC=KEY,ERR=5)AFLO
      KEY=INI+((JSORCE-1)*12)+((KTARG-1)*12*20)
      READ (UNIT=13,FMT='(E9.3)',REC=KEY,ERR=5)AFHI
      DIFF=((AFHI-AFLO)*(E-ELO))/(ENI-ELO)
      INTRPT=AFLO+DIFF
      RETURN
5 CALL CLEAR
      WRITE (*,10)
10 FORMAT(///,' ERROR: Unable to read the value of absorbed fraction f
&or the given photon      energy, source, and target from file "AB
&SFRAC"',/, ' SOURCE: INTRPT FORTRAN',/, ' CORRECTIVE ACTION: Check t
&he nuclide decay scheme and try again! ',///)
      ICHECK=1
      RETURN
      END
```

PCLASS.FOR

10-15-1987

```

*****
*
*      SUBROUTINE NAME : PCLASS FORTRAN
*      PURPOSE: To explain the pulmonary clearance classification
*              of inorganic compounds
*
*****

SUBROUTINE PCLASS (IVISH)
1 CALL CLEAR
PRINT *, 'PULMONARY CLEARANCE CLASSIFICATION OF INORGANIC COMPODS.'
PRINT *, '-----'
PRINT *
PRINT *, 'CLASS Y--> AVIO RETENTION: CLEARED SLOWLY (YEARS) '
PRINT *, '-----'
PRINT *, ' Carbides--> actinides,lanthanides,Zr,Y,Mn '
PRINT *, ' Sulfides--> none '
PRINT *, ' Sulfates--> none '
PRINT *, ' Carbonates--> none '
PRINT *, ' Phosphates--> none '
PRINT *, ' Oxides and hydroxides--> lanthanides, actinide Groups '
PRINT *, ' 8 (V and VI), 1b,2b(IV and V) '
PRINT *, ' 3b except Sc(3+),and 6b '
PRINT *, ' Halides--> lanthanide fluorides '
PRINT *, ' Nitrates--> none '
PRINT *
PAUSE ' TO RESUME PRESS <RETURN>!!'
CALL CLEAR
PRINT *, 'Class W--> Moderate retention: intermediate rate (weeks)'
PRINT *, '-----'
PRINT *, ' Carbides--> Cations of all Class W hydroxides except '
PRINT *, ' those listed as Class Y carbides '
PRINT *, ' Sulfides--> Groups 2a(V + VI), 4a(IV-VI), 5a(IV-VI),1b '
PRINT *, ' 2b, and 6b(V+VI) '
PRINT *, ' Sulfates--> Groups 2c(IV-VII), and 5a(IV-VI) '
PRINT *, ' Carbonates--> lanthanides, Bi(3+), Group 2c(IV-VII) '
PRINT *, ' Phosphates--> Zn(2+), Sn(3+), Mg(2+), Fe(3+), Bi(3+), '
PRINT *, ' and lanthanides '
PRINT *, ' Oxides and hydroxides--> Groups 2c(II-VII),3c(III-VI), '
PRINT *, ' 4a(III-VI), 5a(IV-VI), 6a(IV-VI), '
PRINT *, ' 1b, 8, 2b(VI),4b, 5b, and 7b '
PRINT *, ' Sc(3+) '
PRINT *, ' Halides--> lanthanides (except fluorides), Groups 2a,3a '
PRINT *, ' (III-VI), 4a(IV-VI),5a(IV-VI),8,1b,2b,3b(IV '
PRINT *, ' -V),4b,5b,6b, and 7b '
PRINT *, ' Nitrates--> all cations whose hydroxides are Class Y '
PRINT *, ' and W '
PRINT *
PAUSE ' TO RESUME PRESS <RETURN>!!'
CALL CLEAR
PRINT *, 'Class O--> Minimal retention: rapid clearance (days) '
PRINT *, '-----'
PRINT *, ' Carbides--> all hydroxides '
PRINT *, ' Sulfides--> all except Class W '
PRINT *, ' Sulfates--> all except Class W '

```

PCLASS.FOR

10-15-1987

```
PRINT *, ' Carbonates--> all except Class W          '
```

PRINT *, ' Phosphates--> ell except Class W '

PRINT *, ' Oxides and Hydroxides--> Groups 1a, 3a(II), 4a(II), 5a'

PRINT *, ' (II,III), 6a(III) '

PRINT *, ' Halides--> Groups 1a and 7a '

PRINT *, ' Nitretes--> all except Class W '

PRINT *, ' Noble Gases--> Group 0 '

PRINT *, ' '

PRINT *, 'Note: Where reference is made from one chemical form to '

PRINT *, ' another, it implies that an in vivo convereion '

PRINT *, ' occurs, a.g. hydrolysis reaction '

5 PRINT *

PRINT *, ' Enter integer 1 to repeat or 0 to continue.'

READ (*,*,ERR=5,END=5)NCON

IF (NCON .NE. 1 .AND. (NCON .NE. 0))GOTO 5

IF (NCON .EQ. 1)GOTO 1

CALL CLEAR

END

REFMAN.FOR

10-15-1987

```
*****
*
*      FUNCTION SUBPROGRAM NAME: REFMAN FORTRAN
*      PURPOSE: Provide mass of a target organ in a "reference man"
*
*****

FUNCTION REFMAN(KTARG)
  IF (KTARG .EQ. 1) THEN
    REFMAN=999.
  ELSE IF (KTARG .EQ. 2) THEN
    REFMAN=19.6
  ELSE IF (KTARG .EQ. 3) THEN
    REFMAN=37.1
  ELSE IF (KTARG .EQ. 4) THEN
    REFMAN=8.27
  ELSE IF (KTARG .EQ. 5) THEN
    REFMAN=1500.
  ELSE IF (KTARG .EQ. 6) THEN
    REFMAN=150.
  ELSE IF (KTARG .EQ. 7) THEN
    REFMAN=1040.
  ELSE IF (KTARG .EQ. 8) THEN
    REFMAN=209.
  ELSE IF (KTARG .EQ. 9) THEN
    REFMAN=160.
  ELSE IF (KTARG .EQ. 10) THEN
    REFMAN=1810.
  ELSE IF (KTARG .EQ. 11) THEN
    REFMAN=284.
  ELSE IF (KTARG .EQ. 12) THEN
    REFMAN=45.1
  ELSE IF (KTARG .EQ. 13) THEN
    REFMAN=48200.
  ELSE IF (KTARG .EQ. 14) THEN
    REFMAN=10500.
  ELSE IF (KTARG .EQ. 15) THEN
    REFMAN=2830.
  ELSE IF (KTARG .EQ. 16) THEN
    REFMAN=174.
  ELSE IF (KTARG .EQ. 17) THEN
    REFMAN=65.4
  ELSE IF (KTARG .EQ. 18) THEN
    REFMAN=60.3
  ELSE IF (KTARG .EQ. 19) THEN
    REFMAN=69900.
  END IF
  RETURN
END
```

RESPIR.FOR

10-15-1987

```
*****
*
*      SUBROUTINE NAME : RESPIR FORTRAN
*
*      PURPOSE: Provide fraction of material deposited, and its
*               clearance rate from each compartment of the lung
*
*****
```

```
      SUBROUTINE RESPIR (CLASS,FA,FB,FC,FD,FE,FF,FG,FH,FI,FJ,CLA,CLB,CLC
&,CLD,CLE,CLF,CLG,CLH,CLI,CLJ)
      CHARACTER*1 CLASS
```

```
*****
*
*      DESCRIPTION OF VARIABLES
*      -----
*
```

```
FA--> Fraction of material deposited in the body fluid compart-
*      ment from the nasal passage region (N-P)
*
FB--> Fraction deposited in the GI tract from the nasal passage
*
FC--> Fraction deposited in the body fluid compartment from the
*      trachea and bronchial tree region (T-B)
*
FD--> Fraction deposited in the GI tract from the trachea and
*      bronchial tree region (T-B)
*
FE--> Fraction deposited in the body fluids from the pulmonary
*      region (P)
*
FF--> Fraction transferred to T-B region from P region with a
*      half-life
*
FG--> Fraction transferred to T-B region from P region with a
*      different half-life
*
FH--> Fraction transferred to the lymphatic system (L) from
*      P region
*
FI--> Fraction deposited in the body fluids from L region with
*      a half-life
*
FJ--> Fraction deposited in the body fluids from L region with
*      a different half-life
*
```

```
      CLEARANCE RATES (/DAYS)
      -----
*
CLA--> Clearance rate of material from N-P to body fluids
*
CLB--> Clearance rate of material from N-P to GI tract
*
CLC--> Clearance rate of material from T-B to body fluids
*
CLD--> Clearance rate of material from T-B to GI tract
*
CLE--> Clearance rate of material from P to body fluids
*
CLF--> Clearance rate of material from P to T-B region
*
CLG--> Clearance rate of material from P to T-B region
*
CLH--> Clearance rate of material from P to L region
*
CLI--> Clearance rate of material from L to body fluids
*
CLJ--> Clearance rate of material from L to body fluids
*
```

```
*****
      IF (CLASS .EQ. 'D') THEN
         FA=0.5
         FB=0.5
         FC=0.95
         FD=0.05
         FE=0.8

```

RESPIR.FOR

10-15-1987

```
FH=0.2
FI=1.0
CLA=LOG(2.)/0.01
CLB=LOG(2.)/0.01
CLC=LOG(2.)/0.01
CLO=LOG(2.)/0.2
CLE=LOG(2.)/0.5
CLH=LOG(2.)/0.5
CLI=LOG(2.)/0.5
ELSE IF (CLASS .EQ. 'W') THEN
  FA=0.1
  FB=0.9
  FC=0.5
  FO=0.5
  FE=0.15
  FF=0.4
  FG=0.4
  FH=0.05
  FI=1.0
  CLA=LOG(2.)/0.01
  CLB=LOG(2.)/0.40
  CLC=LOG(2.)/0.01
  CLO=LOG(2.)/0.2
  CLE=LOG(2.)/50.
  CLF=LOG(2.)/1.0
  CLG=LOG(2.)/50.
  CLH=LOG(2.)/50.
  CLI=LOG(2.)/50.
ELSE IF (CLASS .EQ. 'Y') THEN
  FA=0.01
  FB=0.99
  FC=0.01
  FO=0.99
  FE=0.05
  FF=0.4
  FG=0.4
  FH=0.15
  FI=0.9
  FJ=0.1
  CLA=LOG(2.)/0.01
  CLB=LOG(2.)/0.40
  CLC=LOG(2.)/0.01
  CLO=LOG(2.)/0.2
  CLE=LOG(2.)/500.
  CLF=LOG(2.)/1.0
  CLG=LOG(2.)/500.
  CLH=LOG(2.)/500.
  CLI=LOG(2.)/1000.
  CLJ=0.
END IF
END
```

RESULT.FOR

10-15-1987

```

*****
*
*      SUBROUTINE NAME : RESULT FORTRAN
*
*      PURPOSE: Evaluate Weighted Committed Dose Equivalent using
*               the 10% exclusion principle, the DAC, and the ALI
*
*      DATA FILE REQUIRED :  e) RETENT FILE
*
*****
      SUBROUTINE RESULT (NFIFTY, WDOSE, ALI, POST, IRGANT, DAC, KZ, REMDR, WREMD
&R, WTF, SUM)
      DIMENSION NFIFTY(1:24), WDOSE(1:24), TEMP(1:12), REM(1:5), NTARG(1:5)
      CHARACTER*20 C2
      REAL MAXDOS
      OPEN (UNIT=15, FILE='RETENT', ACCESS='DIRECT', FORM='FORMATTED', RECL=
&66, STATUS='OLD')
*****
*
*      Finding the five organs or tissues of the remainder receiving
*      the highest dose equivalent; the exposure of all other
*      remaining tissues is neglected
*
*****
      DO 1 I=1,20
      WDOSE(I)=0.
1  CONTINUE
      DO 5 J=1,12
      IF (I .GE. 8) THEN
          TEMP(I)=NFIFTY(I+7)
      ELSE
          TEMP(I)=NFIFTY(I+5)
      END IF
5  CONTINUE
      DO 10 I=1,5
      REM(I)=AMAX1(TEMP(1),TEMP(2),TEMP(3),TEMP(4),TEMP(5),TEMP(6),TEMP(
&7),TEMP(8),TEMP(9),TEMP(10),TEMP(11),TEMP(12))
      DO 15 J=1,12
      IF (REM(I) .EQ. TEMP(J)) THEN
          TEMP(J)=0.
          GOTD 10
      END IF
15  CONTINUE
10  CONTINUE
*****
*
*      Weighted committed dose equivalent, WDOSE
*
*****
      DO 20 I=1,19
      IF (I .EQ. 1) THEN
          WT=0.12
          WDOSE(I)=WT*NFIFTY(I)
      ELSE IF (I .EQ. 2) THEN
          WT=0.03
          WDOSE(I)=WT*NFIFTY(I)

```

RESULT. FOR

10-15-1987

```

ELSE IF(I .EQ. 3 .OR. (I .EQ. 4))THEN
  WT=0.25
  WDOSE(I)=WT*HFIFTY(I)
ELSE IF(I .EQ. 5)THEN
  WT=0.12
  WDOSE(I)=WT*HFIFTY(I)
ELSE IF(I .EQ. 13)THEN
  WT=0.15
  WDOSE(I)=WT*HFIFTY(I)
ELSE IF (I .EQ. 14)THEN
  WT=0.03
  WDOSE(I)=WT*HFIFTY(I)
ELSE
  WT=0.06
  DO 25 J=1,5
    IF (HFIFTY(I) .EQ. REM(J))THEN
      WDOSE(I)=WT*REM(J)
      GOTO 20
    END IF
  25 CONTINUE
  END IF
20 CONTINUE
*****
*
*   The maximum weighted committed dose equivalent, MAXDOS
*
*****
  MAXDOS=AMAX1(WDOSE(1),WDOSE(2),WDOSE(3),WDOSE(4),WDOSE(5),WDOSE(6)
&,WDOSE(7),WDOSE(8),WDOSE(9),WDOSE(10),WDOSE(11),WDOSE(12),WDOSE(13)
&,WDOSE(14),WDOSE(15),WDOSE(16),WDOSE(17),WDOSE(18),WDOSE(19))
*****
*
*   Weighted committed dose equivalent,WDOSE which is greater
*   than or equal to 10% of the maximum weighted value of H-50
*   per unit intake in any tissue,MAXDOS
*
*****
  PERC=0.1*MAXDOS
  DO 30 I=1,19
    IF (WDOSE(I) .LT. PERC)THEN
      WDOSE(I)=0.
    END IF
  30 CONTINUE
*****
*
*   Check for the organ named in the metabolic model
*
*****
  ISAVE=(5*(KZ-1))
  SUM=0.
  DO 35 I=1,5
    NTARG(I)=0
  35 CONTINUE
  DO 40 I=1,5

```

RESULT.FOR

10-15-1987

```
KEY=ISAVE+1
READ(UNIT=15,FMT='(A20)',REC=KEY)C2
ISORCE=11(C2)
IF(ISORCE.EQ.6)THEN
  NTARG(1)=11
ELSE IF(ISORCE.EQ.7)THEN
  NTARG(2)=10
ELSE IF(ISORCE.EQ.11)THEN
  NTARG(3)=18
ELSE IF(ISORCE.EQ.14)THEN
  NTARG(4)=16
ELSE IF(ISORCE.EQ.17)THEN
  NTARG(5)=19
ELSE
  GOTO 40
END IF
40 CONTINUE
*****
*
*   Assigning a committed dose equivalent to the "REMAINDER" with
*   weighting factor
*
*****
DO 45 I=1,19
  IF (WDOSE(I).EQ.0.)THEN
    HFIFTY(I)=0.
  END IF
45 CONTINUE
DO 50 I=1,5
  REM(I)=0.
50 CONTINUE
ICOUNT=0
DO 55 I=10,19
  IF (I.EQ.13.OR.(I.EQ.14))GOTO 55
DO 60 J=1,5
  IF (I.EQ. NTARG(J))GOTO 55
60 CONTINUE
  IF (HFIFTY(I).GT.0.)THEN
    ICOUNT=ICOUNT+1
    REM(ICOUNT)=HFIFTY(I)
    HFIFTY(I)=0.
    WDOSE(I)=0.
  END IF
55 CONTINUE
REMR=AMAX1(REM(1),REM(2),REM(3),REM(4),REM(5))
WTF=0.06*ICOUNT
WREMR=WTF*REMR
*****
*
*   Annual limit on intake, ALI
*
*****
SUM=WREMR
DO 65 I=1,19
```

RESULT.FOR

10-15-1987

```
IF (WOOSE(I) .EQ. 0.)GOTO 65
SUM=SUM+WOOSE(I)
65 CONTINUE
POST=0.05/SUM
ALI=POST
SAVE=0.
IRGANT=0
DO 70 I=1,19
IF (HFIFTY(I) .EQ. 0.)GOTO 70
POST1=0.5/POST
IF (HFIFTY(I) .GT. POST1)THEN
  IF (HFIFTY(I) .GT. SAVE)THEN
    SAVE=HFIFTY(I)
    IRGANT=I
  ELSE
    GOTO 70
  END IF
END IF
END IF
70 CONTINUE
IF (SAVE .NE. 0.)THEN
  ALI=0.5/SAVE
END IF
```

```
*****
*
*   Derived air concentration, DAC
*
*****
```

```
DAC=ALI/2.4E+03
RETURN
END
```

SOURCE.FOR

10-15-1987

```
*****
*
*      FUNCTION SUBPROGRAM NAME: SOURCE FORTRAN
*
*      PURPOSE: Provide mass of the source organ when source organ
*               integer is given as input
*
*****

      FUNCTION SOURCE(ICOMP)
      IF (ICOMP .EQ. 6) THEN
        SOURCE=310.
      ELSE IF (ICOMP .EQ. 7) THEN
        SOURCE=1800.
      ELSE IF (ICOMP .EQ. 10) THEN
        SOURCE=11.
      ELSE IF (ICOMP .EQ. 11) THEN
        SOURCE=100.
      ELSE IF (ICOMP .EQ. 12) THEN
        SOURCE=5000.
      ELSE IF (ICOMP .EQ. 14) THEN
        SOURCE=180.
      ELSE IF (ICOMP .EQ. 15) THEN
        SOURCE=35.
      ELSE IF (ICOMP .EQ. 16) THEN
        SOURCE=20.
      ELSE IF (ICOMP .EQ. 17) THEN
        SOURCE=70000.
      ELSE IF (ICOMP .EQ. 19) THEN
        SOURCE=1450.
      ELSE IF (ICOMP .EQ. 20) THEN
        SOURCE=14.
      ELSE IF (ICOMP .EQ. 21) THEN
        SOURCE=1500.
      END IF
      RETURN
      END
```


SPEFF.FOR

10-15-1987

```
*****
*
* FUNCTION SUBPROGRAM NAME : SPEFF FORTRAN
*
* PURPOSE: Calculates the specific effective energy deposited
*          in target organ, KTARG due to source organ, JSORCE
*
* AUXILIARY PROGRAM REQUIRED: a) DECAY FORTRAN
*                           b) INTRPT FORTRAN
*                           c) ENERGY FORTRAN
*
*****
FUNCTION SPEFF(WORD,JSORCE,KTARG,TMASS,LOOP,MOTS,PLIFE)
REAL INTRPT
DIMENSION EALPHA(1:20),YALPHA(1:20),EBETA(1:50),YBETA(1:50),EPOST(
&1:15),YPOST(1:15),EELEC(1:115),YELEC(1:115),EGAMMA(1:190),YGAMMA(1
&:190)
COMMON EALPHA,YALPHA,EBETA,YBETA,EPOST,YPOST,EELEC,YELEC,EGAMMA,YG
&AMMA,M,11,13,15,17,HLIFE
CHARACTER*8 SAVE,WORD,MOTS
*****
*
* Comparing source organ, JSORCE end target, KTARG. When
* source organ is not equal to the target organ, ICOM=1
* else ICOM=0
*
*****
IF(JSORCE .EQ. 1 .AND. KTARG .EQ. 12)THEN
  ICOM=0
ELSE IF(JSORCE .EQ. 2 .AND. KTARG .EQ. 6)THEN
  ICOM=0
ELSE IF(JSORCE .EQ. 3 .AND. KTARG .EQ. 7)THEN
  ICOM=0
ELSE IF(JSORCE .EQ. 4 .AND. KTARG .EQ. 8)THEN
  ICOM=0
ELSE IF(JSORCE .EQ. 5 .AND. KTARG .EQ. 9)THEN
  ICOM=0
ELSE IF(JSORCE .EQ. 6 .AND. KTARG .EQ. 11)THEN
  ICOM=0
ELSE IF(JSORCE .EQ. 7 .AND. KTARG .EQ. 10)THEN
  ICOM=0
ELSE IF(JSORCE .EQ. 8 .AND. KTARG .EQ. 1)THEN
  ICOM=0
ELSE IF(JSORCE .EQ. 9 .AND. KTARG .EQ. 13)THEN
  ICOM=0
ELSE IF(JSORCE .EQ. 10 .AND. KTARG .EQ. 4)THEN
  ICOM=0
ELSE IF(JSORCE .EQ. 11 .AND. KTARG .EQ. 18)THEN
  ICOM=0
ELSE IF(JSORCE .EQ. 12 .AND. KTARG .EQ. 5)THEN
  ICOM=0
ELSE IF(JSORCE .EQ. 12 .AND. KTARG .EQ. 14)THEN
  ICOM=0
ELSE IF(JSORCE .EQ. 13 .AND. KTARG .EQ. 15)THEN
  ICOM=0
ELSE IF(JSORCE .EQ. 14 .AND. KTARG .EQ. 16)THEN
```

SPEFF.FOR

10-15-1987

```

      ICOM=0
      ELSE IF(JSORCE .EQ. 15 .AND. KTARG .EQ. 3)THEN
        ICOM=0
      ELSE IF(JSORCE .EQ. 16 .AND. KTARG .EQ. 2)THEN
        ICOM=0
      ELSE IF(JSORCE .EQ. 17 .AND. KTARG .EQ. 19)THEN
        ICOM=0
      ELSE
        ICOM=1
      END IF
*****
*
*      Begin with 'Total body' ss sources.
*
*      The character variable 'SAVE' stores the name of the
*      previous radionuclide and hence when it is equal to
*      'WORD', the function 'SPEFF' does not have to call
*      'DECAY' again
*
*****
      IF (SAVE .EQ. WORD)GOTO 5
      ICOUNT=0
      CALL DECAY (WORD,ICOUNT)
      IF (ICOUNT .EQ. 1)THEN
        WORD(1:5)='SORRY'
        RETURN
      END IF
      5 SPEFF=0.
      ICHECK=0
*****
*      Annihilation photons
*
*****
      IF (I3 .NE. 0)THEN
        DO 6 I=17*1,17*13
          EGAMMA(I)=0.511
          YGAMMA(I)=2.*YPOST(I-17)
        6 CONTINUE
      END IF
*****
      IF(KTARG .EQ. 6 .OR. (KTARG .EQ. 7) .OR. (KTARG .EQ. 8) .OR. (KTAR
&G .EQ. 9) .OR. (KTARG .EQ. 12))THEN
        GOTQ 55
      ELSE IF(KTARG .EQ. 5 .OR. KTARG .EQ. 14)THEN
        GOTQ 105
      END IF
*****
*
*      Organs other than bladder, GI tract and bone
*
*
*      ALPHA
*
*****
      IF (M .EQ. 0)GOTQ 15
      OF=20
      DO 10 I=1,M

```

SPEFF.FOR

10-15-1987

```
IF (ICOM .EQ. 1) THEN
  IF (JSORCE .EQ. 17) THEN
    AF=1./69900.
    SPEFF=SPEFF+(YALPHA(1)*EALPHA(1)*AF*QF)
  ELSE IF (KTARG .EQ. 19) THEN
    IF (JSORCE .EQ. 1) THEN
      AF=45.1/(2*200.*69900.)
    ELSE IF (JSORCE .EQ. 2) THEN
      AF=150./(2*250.*69900.)
    ELSE IF (JSORCE .EQ. 3) THEN
      AF=640./(2*400.*69900.)
    ELSE IF (JSORCE .EQ. 4) THEN
      AF=210./(2*220.*69900.)
    ELSE IF (JSORCE .EQ. 5) THEN
      AF=160./(2*135.*69900.)
    ELSE
      AF=1./69900.
    END IF
    SPEFF=SPEFF+(YALPHA(1)*EALPHA(1)*AF*QF)
  ELSE
    AF=0.
  END IF
ELSE
  AF=1.0
  SPEFF=SPEFF+((YALPHA(1)*EALPHA(1)*AF*QF)/TMASS)
END IF
10 CONTINUE
*****
*
*           BETA
*
*****
15 IF (I1 .EQ. 0) GOTO 25
QF=1
QO 20 I=1,11
IF (ICOM .EQ. 1) THEN
  IF (JSORCE .EQ. 17) THEN
    AF=1./69900.
    SPEFF=SPEFF+(YBETA(1)*EBETA(1)*QF*AF)
  ELSE IF (KTARG .EQ. 19) THEN
    IF (JSORCE .EQ. 1) THEN
      AF=45.1/(2*200.*69900.)
    ELSE IF (JSORCE .EQ. 2) THEN
      AF=150./(2*250.*69900.)
    ELSE IF (JSORCE .EQ. 3) THEN
      AF=640./(2*400.*69900.)
    ELSE IF (JSORCE .EQ. 4) THEN
      AF=210./(2*220.*69900.)
    ELSE IF (JSORCE .EQ. 5) THEN
      AF=160./(2*135.*69900.)
    ELSE
      AF=1./69900.
    END IF
    SPEFF=SPEFF+(YBETA(1)*EBETA(1)*QF*AF)
```

SPEFF.FOR

10-15-1987

```

      ELSE
        AF=0.
      END IF
    ELSE
      AF=1.0
      SPEFF=SPEFF+((YBETA(1)*EBETA(1)*QF*AF)/TMASS)
    END IF
  20 CONTINUE
*****
*                                     *
*                                     *
*                                     *
*****
  25 IF (I3 .EQ. 0)GOTO 35
  QF=1
  DD 30 I=1,I3
  IF (ICOM .EQ. 1)THEN
    IF (JSORCE .EQ. 17)THEN
      AF=1./69900.
      SPEFF=SPEFF+(YPOST(1)*EPOST(1)*QF*AF)
    ELSE IF (KTARG .EQ. 19)THEN
      IF (JSORCE .EQ. 1)THEN
        AF=45.1/(2*200.*69900.)
      ELSE IF (JSORCE .EQ. 2)THEN
        AF=150./(2*250.*69900.)
      ELSE IF (JSORCE .EQ. 3)THEN
        AF=640./(2*400.*69900.)
      ELSE IF (JSORCE .EQ. 4)THEN
        AF=210./(2*220.*69900.)
      ELSE IF (JSORCE .EQ. 5)THEN
        AF=160./(2*135.*69900.)
      ELSE
        AF=1./69900.
      END IF
      SPEFF=SPEFF+(YPOST(1)*EPOST(1)*QF*AF)
    ELSE
      AF=D.
    END IF
  ELSE
    AF=1.0
    SPEFF=SPEFF+((YPOST(1)*EPOST(1)*QF*AF)/TMASS)
  END IF
  30 CONTINUE
*****
*                                     *
*                                     *
*                                     *
*****
  35 IF (I5 .EQ. 0)GOTO 45
  QF=1
  DD 40 I=1,I5
  IF (ICOM .EQ. 1)THEN
    IF (JSORCE .EQ. 17)THEN
      AF=1./69900.

```

SPEFF.FOR

10-15-1987

```

      SPEFF=SPEFF+(YELEC(1)*EELEC(1)*QF*AF)
    ELSE IF (KTARG .EQ. 19) THEN
      IF (JSORCE .EQ. 1) THEN
        AF=45.1/(2*200.*69900.)
      ELSE IF (JSORCE .EQ. 2) THEN
        AF=150./(2*250.*69900.)
      ELSE IF (JSORCE .EQ. 3) THEN
        AF=640./(2*400.*69900.)
      ELSE IF (JSORCE .EQ. 4) THEN
        AF=210./(2*220.*69900.)
      ELSE IF (JSORCE .EQ. 5) THEN
        AF=160./(2*135.*69900.)
      ELSE
        AF=1./69900.
      END IF
      SPEFF=SPEFF+(YELEC(1)*EELEC(1)*QF*AF)
    ELSE
      AF=0.
    END IF
  ELSE
    AF=1.0
    SPEFF=SPEFF+((YELEC(1)*EELEC(1)*QF*AF)/TNASS)
  END IF
40 CONTINUE
*****
*                                     *
*                                     *
*                                     *
*****
45 IF (I7 .EQ. 0) GOTO 195
QF=1
DO 50 I=1,17
  AF=0.
  IF (EGAMMA(I) .LT. 0.01) THEN
    IF (ICOM .EQ. 1) THEN
      IF (JSORCE .EQ. 17) THEN
        AF=1.0/69900.
        SPEFF=SPEFF+(YGAMMA(I)*EGAMMA(I)*QF*AF)
      ELSE IF (KTARG .EQ. 19) THEN
        IF (JSORCE .EQ. 1) THEN
          AF=45.1/(2*200.*69900.)
        ELSE IF (JSORCE .EQ. 2) THEN
          AF=150./(2*250.*69900.)
        ELSE IF (JSORCE .EQ. 3) THEN
          AF=640./(2*400.*69900.)
        ELSE IF (JSORCE .EQ. 4) THEN
          AF=210./(2*220.*69900.)
        ELSE IF (JSORCE .EQ. 5) THEN
          AF=160./(2*135.*69900.)
        ELSE
          AF=1./69900.
        END IF
        SPEFF=SPEFF+(YGAMMA(I)*EGAMMA(I)*QF*AF)
      ELSE

```

SPEFF.FOR

10-15-1987

```
      AF=0.
    END IF
  ELSE
    AF=1.
    SPEFF=SPEFF+((YGAMMA(I)*EGAMMA(I)*QF*AF)/TMASS)
  END IF
ELSE
  *****
  *
  *   The function INTRPT interpolates the values of absorbed
  *   fraction, AF for the given energy, EGAMMA(I)
  *
  *****
  ICHECK=0
  AF=INTRPT(EGAMMA(I),JSORCE,KTARG,ICHECK)
  IF (ICHECK .EQ. 1)THEN
    WORD(1:5)='SORRY'
    RETURN
  END IF
  SPEFF=SPEFF+((YGAMMA(I)*EGAMMA(I)*QF*AF)/TMASS)
  END IF
50 CONTINUE
  GOTO 195
  *****
  *
  *   Target organs of the GI tract and bladder
  *
  *
  *   ALPHA
  *
  *****
55 IF (N .EQ. 0)GOTO 65
  QF=20
  IF (ICOM .EQ. 1)THEN
    IF (JSORCE .EQ. 17)THEN
      AF=1./69900.
    ELSE
      AF=0.
    END IF
  ELSE
    IF (KTARG .EQ. 6)THEN
      AF=(0.5*0.01)/250.
    ELSE IF (KTARG .EQ. 7)THEN
      AF=(0.5*0.01)/400.
    ELSE IF (KTARG .EQ. 8)THEN
      AF=(0.5*0.01)/220.
    ELSE IF (KTARG .EQ. 9)THEN
      AF=(0.5*0.01)/135.
    ELSE IF (KTARG .EQ. 12)THEN
      AF=(0.5*0.01)/200.
    END IF
  END IF
  IF (AF .EQ. 0.)GOTO 65
  DD 60 I=1,M
  SPEFF=SPEFF+(ALPHA(I)*EALPHA(I)*QF*AF)
```

SPEFF.FOR

10-15-1987

```

60 CONTINUE
*
65 QF=1
  IF (ICOM .EQ. 1) THEN
    IF (JSORCE .EQ. 17) THEN
      AF=1./69900.
    ELSE
      AF=0.
    END IF
  ELSE
    IF (KTARG .EQ. 6) THEN
      AF=(0.5*1.0)/250.
    ELSE IF (KTARG .EQ. 7) THEN
      AF=(0.5*1.0)/400.
    ELSE IF (KTARG .EQ. 8) THEN
      AF=(0.5*1.0)/220.
    ELSE IF (KTARG .EQ. 9) THEN
      AF=(0.5*1.0)/135.
    ELSE IF (KTARG .EQ. 12) THEN
      AF=(0.5*1.0)/200.
    END IF
  END IF
  IF (AF .EQ. 0.) GOTO 95
*****
*
*           BETA
*
*****
  IF (I1 .EQ. 0) GOTO 75
  DO 70 I=1,I1
    SPEFF=SPEFF+(YBETA(I)*EBETA(I)*QF*AF)
  70 CONTINUE
*****
*
*           POSITRON
*
*****
  75 IF (I3 .EQ. 0) GOTO 85
  DO 80 I=1,I3
    SPEFF=SPEFF+(YPOST(I)*EPOST(I)*QF*AF)
  80 CONTINUE
*****
*
*           ELECTRON
*
*****
  85 IF (I5 .EQ. 0) GOTO 95
  DO 90 I=1,I5
    SPEFF=SPEFF+(YELEC(I)*EELEC(I)*QF*AF)
  90 CONTINUE
*****
*
*           PHOTON
*
*****

```

SPEFF.FOR

10-15-1987

```

*****
95 IF (.EQ. 0)GOTO 195
  QF=1
  DO 100 I=1,17
    AF=0.
    IF (EGAMMA(I) .LT. 0.01)THEN
      IF (ICOM .EQ. 1)THEN
        IF (JSORCE .EQ. 17)THEN
          AF=1./69900.
          SPEFF=SPEFF+(YGAMMA(I)*EGAMMA(I)*AF*QF)
        ELSE
          AF=0.
        END IF
      ELSE
        IF (KTARG .EQ. 6)THEN
          AF=1.0/(2.*250.)
        ELSE IF (KTARG .EQ. 7)THEN
          AF=1.0/(2.*400.)
        ELSE IF (KTARG .EQ. 8)THEN
          AF=1.0/(2.*220.)
        ELSE IF (KTARG .EQ. 9)THEN
          AF=1.0/(2.*135.)
        ELSE IF (KTARG .EQ. 12)THEN
          AF=1.0/(2.*200.)
        END IF
        SPEFF=SPEFF+(YGAMMA(I)*EGAMMA(I)*AF*QF)
      END IF
    ELSE
      ICHECK=0
      AF=INTRPT(EGAMMA(I),JSORCE,KTARG,ICHECK)
      IF (ICHECK .EQ. 1)THEN
        WORD(1:5)='SORRY'
        RETURN
      END IF
      SPEFF=SPEFF+((YGAMMA(I)*EGAMMA(I)*AF*QF)/THASS)
    END IF
  100 CONTINUE
  GOTO 195
105 IF (LOOP .EQ. 0. .AND. (JSORCE .EQ. 12))GOTO 185
*****
*
*   Target organs in bone
*
*****
  IF (M .EQ. 0)GOTO 125
  QF=20
  DO 120 I=1,M
    IF (ICOM .EQ. 1)THEN
      IF (JSORCE .EQ. 17)THEN
        AF=1./69900.
        SPEFF=SPEFF+(YALPHA(I)*EALPHA(I)*QF*AF)
      ELSE
        AF=0.
      END IF
    
```


SPEFF.FOR

10-15-1987

```

ELSE
*****
*
*   Bone surface calls
*
*****
      IF (KTARG.EQ. 14)THEN
*****
*
*   Trabecular bone
*
*****
      IF (LOOP.EQ. 1)THEN
*****
*   Alpha emitter uniform in volume
*
*****
      IF (MOTS(1:4).EQ. 'P-33'.OR. (MOTS(1:6).EQ. 'NB-93M').OR.
&R. (MOTS(1:5).EQ. 'NB-94').OR. (MOTS(1:5).EQ. 'U-232').OR. (MO
&TS(1:5).EQ. 'U-233').OR. (MOTS(1:5).EQ. 'U-234').OR. (MOTS(1:5
&) .EQ. 'U-235').OR. (MOTS(1:5).EQ. 'U-236').OR. (MOTS(1:5).EQ.
& 'U-238').OR. (MOTS(1:2).EQ. 'NA').OR. (MOTS(1:2).EQ. 'CR').OR.
&R. (MOTS(1:2).EQ. 'RB').OR. (MOTS(1:5).EQ. 'ZN-65').OR. (MOTS(
&1:6).EQ. 'PB-205').OR. (MOTS(1:6).EQ. 'PB-210').OR. (MOTS(1:4)
&.EQ. 'BE-7').OR. (MOTS(1:5).EQ. 'BE-10').OR. (MOTS(1:4).EQ. '
&V-49').OR. (MOTS(1:6).EQ. 'PD-103').OR. (MOTS(1:6).EQ. 'PD-107
&').OR. (MOTS(1:6).EQ. 'SN-113').OR. (MOTS(1:7).EQ. 'SN-119M')
&.OR. (MOTS(1:6).EQ. 'SN-123').OR. (MOTS(1:6).EQ. 'SN-126').OR.
& (MOTS(1:6).EQ. 'TA-182').OR. (MOTS(1:5).EQ. 'W-181').OR. (MOT
&S(1:5).EQ. 'W-185').OR. (MOTS(1:5).EQ. 'W-188'))THEN
      AF=0.025
*****
*   Alkaline earths
*
*****
      ELSE IF (MOTS(1:2).EQ. 'SR'.OR. (MOTS(1:2).EQ. 'CA').OR
&. (MOTS(1:2).EQ. 'RA').OR. (MOTS(1:2).EQ. 'BA'))THEN
      IF (PLIFE.GT. 15)THEN
      AF=0.025
      ELSE
      AF=0.25
      END IF
      ELSE
*****
*   Alpha emitter on bone surfaces
*
*****
      AF=0.25
      END IF
*****
*
*   Cortical bone
*
*****
      ELSE IF (LOOP.GT. 1)THEN
*****
*   Alpha emitter uniform in volume
*

```

SPEFF.FOR

10-15-1987

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*****
      IF (MOTS(1:4) .EQ. 'P-33' .OR. (MOTS(1:6) .EQ. 'NB-93M') .O
&R. (MOTS(1:5) .EQ. 'NB-94') .OR. (MOTS(1:5) .EQ. 'U-232') .OR. (M
&TS(1:5) .EQ. 'U-233') .OR. (MOTS(1:5) .EQ. 'U-234') .OR. (MOTS(1:5
&) .EQ. 'U-235') .OR. (MOTS(1:5) .EQ. 'U-236') .OR. (MOTS(1:5) .EQ.
& 'U-238') .OR. (MOTS(1:2) .EQ. 'NA') .OR. (MOTS(1:2) .EQ. 'CR') .O
&R. (MOTS(1:2) .EQ. 'RB') .OR. (MOTS(1:5) .EQ. 'ZN-65') .OR. (MOTS(
&1:6) .EQ. 'PB-205') .OR. (MOTS(1:6) .EQ. 'PB-210') .OR. (MOTS(1:4)
& .EQ. 'BE-7') .OR. (MOTS(1:5) .EQ. 'BE-10') .OR. (MOTS(1:4) .EQ. '
&V-49') .OR. (MOTS(1:6) .EQ. 'PD-103') .OR. (MOTS(1:6) .EQ. 'PD-107
&') .OR. (MOTS(1:6) .EQ. 'SN-113') .OR. (MOTS(1:7) .EQ. 'SW-119M')
&.OR. (MOTS(1:6) .EQ. 'SN-123') .OR. (MOTS(1:6) .EQ. 'SW-126') .OR.
& (MOTS(1:6) .EQ. 'TA-182') .OR. (MOTS(1:5) .EQ. 'W-181') .OR. (MOT
&S(1:5) .EQ. 'W-185') .OR. (MOTS(1:5) .EQ. 'W-188'))THEN
      AF=0.01
*****
*      Alkaline earths      *
*****
      ELSE IF (MOTS(1:2) .EQ. 'SR' .OR. (MOTS(1:2) .EQ. 'CA') .OR
&. (MOTS(1:2) .EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'BA'))THEN
      IF (PLIFE .GT. 15)THEN
      AF=0.01
      ELSE
      AF=0.25
      END IF
      ELSE
*****
*      Alpha emitter on bone surfaces      *
*****
      AF=0.25
      END IF
      END IF
      SPEFF=SPEFF+((YALPHA(1)*EALPHA(1)*QF*AF)/120.)
*****
*      Red marrow      *
*****
      ELSE IF (KTARG .EQ. 5)THEN
*****
*      Trabecular bone      *
*****
      IF (LOOP .EQ. 1)THEN
*****
*      Alpha emitter uniform in volume      *
*****
      IF (MOTS(1:4) .EQ. 'P-33' .OR. (MOTS(1:6) .EQ. 'NB-93M') .O
&R. (MOTS(1:5) .EQ. 'NB-94') .OR. (MOTS(1:5) .EQ. 'U-232') .OR. (M
&TS(1:5) .EQ. 'U-233') .OR. (MOTS(1:5) .EQ. 'U-234') .OR. (MOTS(1:5
&) .EQ. 'U-235') .OR. (MOTS(1:5) .EQ. 'U-236') .OR. (MOTS(1:5) .EQ.
& 'U-238') .OR. (MOTS(1:2) .EQ. 'NA') .OR. (MOTS(1:2) .EQ. 'CR') .O
&R. (MOTS(1:2) .EQ. 'RB') .OR. (MOTS(1:5) .EQ. 'ZN-65') .OR. (MOTS(

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SPEFF.FOR

10-15-1987

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&1:6) .EQ. 'PB-205') .OR. (MOTS(1:6) .EQ. 'PB-210') .OR. (MOTS(1:4)
& .EQ. 'BE-7') .OR. (MOTS(1:5) .EQ. 'BE-10') .OR. (MOTS(1:4) .EQ. '
&V-49') .OR. (MOTS(1:6) .EQ. 'PD-103') .OR. (MOTS(1:6) .EQ. 'PD-107
&') .OR. (MOTS(1:6) .EQ. 'SN-113') .OR. (MOTS(1:7) .EQ. 'SN-119W')
&.OR. (MOTS(1:6) .EQ. 'SN-123') .OR. (MOTS(1:6) .EQ. 'SN-126') .OR.
& (MOTS(1:6) .EQ. 'TA-182') .OR. (MOTS(1:5) .EQ. 'W-181') .OR. (MOT
&S(1:5) .EQ. 'W-185') .OR. (MOTS(1:5) .EQ. 'W-188')) THEN
    AF=0.05
*****
*      Alkaline earths      *
*****
    ELSE IF (MOTS(1:2) .EQ. 'SR' .OR. (MOTS(1:2) .EQ. 'CA') .OR
& (MOTS(1:2) .EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'BA')) THEN
        IF (PLIFE .GT. 15) THEN
            AF=0.05
        ELSE
            AF=0.5
        END IF
    ELSE
*****
*      Alpha emitter on bone surfaces      *
*****
        AF=0.5
    END IF
    SPEFF=SPEFF+((YALPNA(I)*EALPNA(I)*QF*AF)/1500.)
*****
*      Cortical bone      *
*****
    ELSE IF (LOOP .GT. 1) THEN
        AF=0.0
    END IF
    END IF
    END IF
120 CONTINUE
*
125 QF=1
*****
*      Beta      *
*****
    IF (I1 .EQ. 0) GOTO 165
    DO 160 I=1,I1
    IF (ICOM .EQ. 1) THEN
        IF (JSORCE .EQ. 17) THEN
            AF=1./69900.
            SPEFF=SPEFF+(YBETA(I)*EBETA(I)*QF*AF)
        ELSE
            AF=0.
        END IF
    ELSE
*****

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SPEFF.FOR

10-15-1987

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*
*   Bone surface cells
*
*
*****
      IF (KTARG .EQ. 14) THEN
*****
*
*   Trabecular bone
*
*
*****
      IF (LOOP .EQ. 1) THEN
*****
*   Beta emitter uniform in volume
*
*****
      IF (MOTS(1:4) .EQ. 'P-33' .OR. (MOTS(1:6) .EQ. 'NB-93M') .Q
&R. (MOTS(1:5) .EQ. 'NB-94') .OR. (MOTS(1:5) .EQ. 'U-232') .OR. (MO
&TS(1:5) .EQ. 'U-233') .OR. (MOTS(1:5) .EQ. 'U-234') .OR. (MOTS(1:5
&) .EQ. 'U-235') .OR. (MOTS(1:5) .EQ. 'U-236') .OR. (MOTS(1:5) .EQ.
& 'U-238') .OR. (MOTS(1:2) .EQ. 'NA') .OR. (MOTS(1:2) .EQ. 'CR') .Q
&R. (MOTS(1:2) .EQ. 'RB') .OR. (MOTS(1:5) .EQ. 'ZN-65') .OR. (MOTS(
&1:6) .EQ. 'PB-205') .OR. (MOTS(1:6) .EQ. 'PB-210') .OR. (MOTS(1:4)
& .EQ. 'BE-7') .OR. (MOTS(1:5) .EQ. 'BE-10') .OR. (MOTS(1:4) .EQ. '
&V-49') .OR. (MOTS(1:6) .EQ. 'PD-103') .OR. (MOTS(1:6) .EQ. 'PD-107
&') .OR. (MOTS(1:6) .EQ. 'SN-113') .OR. (MOTS(1:7) .EQ. 'SN-119M')
&.OR. (MOTS(1:6) .EQ. 'SN-123') .OR. (MOTS(1:6) .EQ. 'SN-126') .OR.
& (MOTS(1:6) .EQ. 'TA-182') .OR. (MOTS(1:5) .EQ. 'U-181') .OR. (MOT
&S(1:5) .EQ. 'U-185') .OR. (MOTS(1:5) .EQ. 'U-188')) THEN
          AF=0.025
*****
*   Alkaline earths
*
*****
      ELSE IF (MOTS(1:2) .EQ. 'SR' .OR. (MOTS(1:2) .EQ. 'CA') .OR
&. (MOTS(1:2) .EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'BA')) THEN
          IF (PLIFE .GT. 15) THEN
              AF=0.025
          ELSE
              IF (EBETA(1) .LT. 0.2) THEN
                  AF=0.25
              ELSE
                  AF=0.025
              END IF
          END IF
      ELSE
*****
*   Beta emitter on bone surfaces
*
*****
      IF (EBETA(1) .LT. 0.2) THEN
          AF=0.25
      ELSE IF (EBETA(1) .GE. 0.2) THEN
          AF=0.025
      END IF
      END IF
*****
*

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SPEFF.FOR

10-15-1987

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*      Cortical bone
*
*****
      ELSE IF (LOOP .GT. 1) THEN
*****
*      Beta emitter uniform in volume
*
*****
      IF (MOTS(1:4) .EQ. 'P-33' .OR. (MOTS(1:6) .EQ. 'NB-93M') .Q
&R. (MOTS(1:5) .EQ. 'NB-94') .OR. (MOTS(1:5) .EQ. 'U-232') .OR. (MO
&TS(1:5) .EQ. 'U-233') .OR. (MOTS(1:5) .EQ. 'U-234') .OR. (MOTS(1:5
&) .EQ. 'U-235') .OR. (MOTS(1:5) .EQ. 'U-236') .OR. (MOTS(1:5) .EQ.
& 'U-238') .OR. (MOTS(1:2) .EQ. 'NA') .OR. (MOTS(1:2) .EQ. 'CR') .Q
&R. (MOTS(1:2) .EQ. 'RB') .OR. (MOTS(1:5) .EQ. 'ZN-65') .OR. (MOTS(
&1:6) .EQ. 'PB-205') .OR. (MOTS(1:6) .EQ. 'PB-210') .OR. (MOTS(1:4)
& .EQ. 'BE-7') .OR. (MOTS(1:5) .EQ. 'BE-10') .OR. (MOTS(1:4) .EQ. '
&V-49') .OR. (MOTS(1:6) .EQ. 'PO-103') .OR. (MOTS(1:6) .EQ. 'PO-107
&') .OR. (MOTS(1:6) .EQ. 'SN-113') .OR. (MOTS(1:7) .EQ. 'SN-119M')
&.OR. (MOTS(1:6) .EQ. 'SN-123') .OR. (MOTS(1:6) .EQ. 'SN-126') .OR.
& (MOTS(1:6) .EQ. 'TA-182') .OR. (MOTS(1:5) .EQ. 'W-181') .OR. (MOT
&S(1:5) .EQ. 'W-185') .OR. (MOTS(1:5) .EQ. 'W-188')) THEN
      AF=0.015
*****
*      Alkaline earths
*
*****
      ELSE IF (MOTS(1:2) .EQ. 'SR' .OR. (MOTS(1:2) .EQ. 'CA') .OR
&. (MOTS(1:2) .EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'BA')) THEN
      IF (PLIFE .GT. 15) THEN
        AF=0.015
      ELSE
        IF (EBETA(1) .LT. 0.2) THEN
          AF=0.25
        ELSE
          AF=0.015
        END IF
      END IF
    ELSE
*****
*      Beta emitter on bone surfaces
*
*****
      IF (EBETA(1) .LT. 0.2) THEN
        AF=0.25
      ELSE IF (EBETA(1) .GE. 0.2) THEN
        AF=0.015
      END IF
    END IF
  END IF
  SPEFF=SPEFF+((YBETA(1)*EBETA(1)*QF*AF)/120.)
*****
*      Red marrow
*
*****
      ELSE
*****

```

SPEFF.FOR

10-15-1987

```

*
*   Trabecular bone
*
*****
      IF (LOOP.EQ. 1) THEN
*****
*   Beta emitter uniform in volume
*
*****
      IF (MOTS(1:4).EQ. 'P-33'.OR. (MOTS(1:6).EQ. 'NB-93H').OR.
&R. (MOTS(1:5).EQ. 'NB-94').OR. (MOTS(1:5).EQ. 'U-232').OR. (MOTS(1:5).EQ. 'U-233').OR. (MOTS(1:5).EQ. 'U-234').OR. (MOTS(1:5).EQ. 'U-235').OR. (MOTS(1:5).EQ. 'U-236').OR. (MOTS(1:5).EQ. 'U-238').OR. (MOTS(1:2).EQ. 'NA').OR. (MOTS(1:2).EQ. 'CR').OR.
&R. (MOTS(1:2).EQ. 'RB').OR. (MOTS(1:5).EQ. 'ZN-65').OR. (MOTS(1:6).EQ. 'PB-205').OR. (MOTS(1:6).EQ. 'PB-210').OR. (MOTS(1:4).EQ. 'BE-7').OR. (MOTS(1:5).EQ. 'BE-10').OR. (MOTS(1:4).EQ. 'V-49').OR. (MOTS(1:6).EQ. 'PD-103').OR. (MOTS(1:6).EQ. 'PD-107').OR. (MOTS(1:6).EQ. 'SN-113').OR. (MOTS(1:7).EQ. 'SN-119H').OR. (MOTS(1:6).EQ. 'SN-123').OR. (MOTS(1:6).EQ. 'SN-126').OR.
& (MOTS(1:6).EQ. 'TA-182').OR. (MOTS(1:5).EQ. 'U-181').OR. (MOTS(1:5).EQ. 'U-185').OR. (MOTS(1:5).EQ. 'U-188')) THEN
          AF=0.35
*****
*   Alkaline earths
*
*****
      ELSE IF (MOTS(1:2).EQ. 'SR'.OR. (MOTS(1:2).EQ. 'CA').OR.
&. (MOTS(1:2).EQ. 'RA').OR. (MOTS(1:2).EQ. 'BA')) THEN
          IF (PLIFE.GT. 15) THEN
              AF=0.35
          ELSE
              AF=0.5
          END IF
      ELSE
*****
*   Beta emitter on bone surfaces
*
*****
          AF=0.5
      END IF
      SPEFF=SPEFF+((YBETA(I)*EBETA(I)*AF)/1500.)
*****
*
*   Cortical bone
*
*****
      ELSE IF (LOOP.GT. 1) THEN
          AF=0.0
      END IF
      END IF
      END IF
160 CONTINUE
*****
*
*   Positron
*
*****

```

SPEFF.FOR

10-15-1987

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*****
165 IF (I3 .EQ. 0)GOTO 175
    DO 170 I=1,I3
    IF (ICOM .EQ. 1)THEN
        IF (JSORCE .EQ. 17)THEN
            AF=1./69900.
            SPEFF=SPEFF+(YPOST(I)*EPOST(I)*QF*AF)
        ELSE
            AF=0.
        END IF
    ELSE
*****
*
*   Bone surface cells
*
*****
        IF (KTARG .EQ. 14)THEN
*****
*
*   Trabecular bone
*
*****
            IF (LOOP .EQ. 1)THEN
*****
*   Positron emitter uniform in volume
*****
                IF (MOTS(1:4) .EQ. 'P-33' .OR. (MOTS(1:6) .EQ. 'NB-93M') .O
&R. (MOTS(1:5) .EQ. 'NB-94') .OR. (MOTS(1:5) .EQ. 'U-232') .OR. (MO
&TS(1:5) .EQ. 'U-233') .OR. (MOTS(1:5) .EQ. 'U-234') .OR. (MOTS(1:5
&) .EQ. 'U-235') .OR. (MOTS(1:5) .EQ. 'U-236') .OR. (MOTS(1:5) .EQ.
& 'U-238') .OR. (MOTS(1:2) .EQ. 'NA') .OR. (MOTS(1:2) .EQ. 'CR') .O
&R. (MOTS(1:2) .EQ. 'RB') .OR. (MOTS(1:5) .EQ. 'ZN-65') .OR. (MOTS(
&1:6) .EQ. 'PB-205') .OR. (MOTS(1:6) .EQ. 'PB-210') .OR. (MOTS(1:4)
& .EQ. 'BE-7') .OR. (MOTS(1:5) .EQ. 'BE-10') .OR. (MOTS(1:4) .EQ. '
&V-49') .OR. (MOTS(1:6) .EQ. 'PD-103') .OR. (MOTS(1:6) .EQ. 'PD-107
&') .OR. (MOTS(1:6) .EQ. 'SN-113') .OR. (MOTS(1:7) .EQ. 'SN-119M')
&.OR. (MOTS(1:6) .EQ. 'SN-123') .OR. (MOTS(1:6) .EQ. 'SN-126') .OR.
& (MOTS(1:6) .EQ. 'TA-182') .OR. (MOTS(1:5) .EQ. 'W-181') .OR. (MO
&S(1:5) .EQ. 'W-185') .OR. (MOTS(1:5) .EQ. 'W-188'))THEN
                    AF=0.025
*****
*   Alkaline earths
*
*****
                ELSE IF (MOTS(1:2) .EQ. 'SR' .OR. (MOTS(1:2) .EQ. 'CA') .OR
&. (MOTS(1:2) .EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'BA'))THEN
                    IF (PLIFE .GT. 15)THEN
                        AF=0.025
                    ELSE
                        IF (EPOST(I) .LT. 0.2)THEN
                            AF=0.25
                        ELSE
                            AF=0.025
                        END IF
                    END IF
                END IF
            END IF
        END IF
    END IF

```

SPEFF.FOR

10-15-1987

```

      ELSE
*****
*      Positron emitter on bone surfaces      *
*****
      IF (EPOST(1) .LT. 0.2) THEN
        AF=0.25
      ELSE
        AF=0.025
      END IF
    END IF
*****
*
*      Cortical bone
*
*****
      ELSE IF (LOOP .GT. 1) THEN
*****
*      Positron emitter uniform in volume      *
*****
      IF (MOTS(1:4) .EQ. 'P-33' .OR. (MOTS(1:6) .EQ. 'NB-93M') .OR.
        &R. (MOTS(1:5) .EQ. 'NB-94') .OR. (MOTS(1:5) .EQ. 'U-232') .OR. (MOTS(1:5) .EQ. 'U-233') .OR. (MOTS(1:5) .EQ. 'U-234') .OR. (MOTS(1:5) .EQ. 'U-235') .OR. (MOTS(1:5) .EQ. 'U-236') .OR. (MOTS(1:5) .EQ. 'U-238') .OR. (MOTS(1:2) .EQ. 'NA') .OR. (MOTS(1:2) .EQ. 'CR') .OR.
        &R. (MOTS(1:2) .EQ. 'RB') .OR. (MOTS(1:5) .EQ. 'Zn-65') .OR. (MOTS(1:6) .EQ. 'PB-205') .OR. (MOTS(1:6) .EQ. 'PB-210') .OR. (MOTS(1:4) .EQ. 'BE-7') .OR. (MOTS(1:5) .EQ. 'BE-10') .OR. (MOTS(1:4) .EQ. 'V-49') .OR. (MOTS(1:6) .EQ. 'PO-103') .OR. (MOTS(1:6) .EQ. 'PO-107') .OR. (MOTS(1:6) .EQ. 'SN-113') .OR. (MOTS(1:7) .EQ. 'SN-119M') .OR. (MOTS(1:6) .EQ. 'SN-123') .OR. (MOTS(1:6) .EQ. 'SN-126') .OR.
        & (MOTS(1:6) .EQ. 'TA-182') .OR. (MOTS(1:5) .EQ. 'W-181') .OR. (MOTS(1:5) .EQ. 'W-185') .OR. (MOTS(1:5) .EQ. 'W-188')) THEN
        AF=0.015
*****
*      Alkaline earths
*
*****
      ELSE IF (MOTS(1:2) .EQ. 'SR' .OR. (MOTS(1:2) .EQ. 'CA') .OR.
        & (MOTS(1:2) .EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'BA')) THEN
        IF (PLIFE .GT. 15) THEN
          AF=0.015
        ELSE
          IF (EPOST(1) .LT. 0.2) THEN
            AF=0.25
          ELSE
            AF=0.015
          END IF
        END IF
      ELSE
*****
*      Positron emitter on bone surfaces
*
*****
      IF (EPOST(1) .LT. 0.2) THEN
        AF=0.25
      ELSE

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SPEFF.FOR

10-15-1987

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      AF=0.015
      ENO IF
    END IF
  END IF
  SPEFF=SPEFF+((YPOST(1)*EPOST(1)*QF*AF)/120.)
  *****
  *
  *   Red marrow
  *
  *****
  ELSE
  *****
  *
  *   Trabecular bone
  *
  *****
  IF (LOOP .EQ. 1) THEN
  *****
  *   Positron emitter uniform in volume
  *
  *****
    IF (MOTS(1:4) .EQ. 'P-33' .OR. (MOTS(1:6) .EQ. 'NB-93H') .OR.
      & (MOTS(1:5) .EQ. 'NB-94') .OR. (MOTS(1:5) .EQ. 'U-232') .OR. (MOTS(1:5) .EQ. 'U-233') .OR. (MOTS(1:5) .EQ. 'U-234') .OR. (MOTS(1:5) .EQ. 'U-235') .OR. (MOTS(1:5) .EQ. 'U-236') .OR. (MOTS(1:5) .EQ. 'U-238') .OR. (MOTS(1:2) .EQ. 'NA') .OR. (MOTS(1:2) .EQ. 'CR') .OR. (MOTS(1:2) .EQ. 'RB') .OR. (MOTS(1:5) .EQ. 'ZN-65') .OR. (MOTS(1:6) .EQ. 'PB-205') .OR. (MOTS(1:6) .EQ. 'PB-210') .OR. (MOTS(1:4) .EQ. 'BE-7') .OR. (MOTS(1:5) .EQ. 'BE-10') .OR. (MOTS(1:4) .EQ. 'V-49') .OR. (MOTS(1:6) .EQ. 'PD-103') .OR. (MOTS(1:6) .EQ. 'PD-107') .OR. (MOTS(1:6) .EQ. 'SN-113') .OR. (MOTS(1:7) .EQ. 'SN-119M') .OR. (MOTS(1:6) .EQ. 'SN-123') .OR. (MOTS(1:6) .EQ. 'SN-126') .OR. (MOTS(1:6) .EQ. 'TA-182') .OR. (MOTS(1:5) .EQ. 'U-181') .OR. (MOTS(1:5) .EQ. 'U-185') .OR. (MOTS(1:5) .EQ. 'U-188')) THEN
      AF=0.35
  *****
  *   Alkaline earths
  *
  *****
    ELSE IF (MOTS(1:2) .EQ. 'SR' .OR. (MOTS(1:2) .EQ. 'CA') .OR.
      & (MOTS(1:2) .EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'BA')) THEN
      IF (PLIFE .GT. 15) THEN
        AF=0.35
      ELSE
        AF=0.5
      END IF
    ELSE
  *****
  *   Positron emitter on bone surfaces
  *
  *****
    AF=0.5
  ENO IF
  SPEFF=SPEFF+((YPOST(1)*EPOST(1)*QF*AF)/1500.)
  *****
  *
  *   Cortical bone
  *
  *****

```

SPEFF.FOR

10-15-1987

```

*
*****
      ELSE IF (LOOP .GT. 1) THEN
        AF=0.0
      END IF
    END IF
  END IF
170 CONTINUE
*****
*
*   Electron
*
*****
175 IF (15 .EQ. 0) GOTO 182
DO 180 I=1,15
  IF (ICOM .EQ. 1) THEN
    IF (JSORCE .EQ. 17) THEN
      AF=1./69900.
      SPEFF=SPEFF+(YELEC(I)*EELEC(I)*QF*AF)
    ELSE
      AF=0.
    END IF
  ELSE
*****
*
*   Bone surface cells
*
*****
    IF (KTARG .EQ. 14) THEN
*****
*
*   Trabecular bone
*
*****
    IF (LOOP .EQ. 1) THEN
*****
*   Electron emitter uniform in volume
*
*****
      IF (MOTS(1:4) .EQ. 'P-33' .OR. (MOTS(1:6) .EQ. 'NB-93M') .OR.
        & (MOTS(1:5) .EQ. 'NB-94') .OR. (MOTS(1:5) .EQ. 'U-232') .OR. (MOTS(1:5) .EQ. 'U-233') .OR. (MOTS(1:5) .EQ. 'U-234') .OR. (MOTS(1:5) .EQ. 'U-235') .OR. (MOTS(1:5) .EQ. 'U-236') .OR. (MOTS(1:5) .EQ. 'U-238') .OR. (MOTS(1:2) .EQ. 'NA') .OR. (MOTS(1:2) .EQ. 'CR') .OR.
        & (MOTS(1:2) .EQ. 'RB') .OR. (MOTS(1:5) .EQ. 'ZN-65') .OR. (MOTS(1:6) .EQ. 'PB-205') .OR. (MOTS(1:6) .EQ. 'PB-210') .OR. (MOTS(1:4) .EQ. 'BE-7') .OR. (MOTS(1:5) .EQ. 'BE-10') .OR. (MOTS(1:4) .EQ. 'Bv-49') .OR. (MOTS(1:6) .EQ. 'PD-103') .OR. (MOTS(1:6) .EQ. 'PD-107') .OR. (MOTS(1:6) .EQ. 'SN-113') .OR. (MOTS(1:7) .EQ. 'SN-119M') .OR. (MOTS(1:6) .EQ. 'SN-123') .OR. (MOTS(1:6) .EQ. 'SN-126') .OR.
        & (MOTS(1:6) .EQ. 'TA-182') .OR. (MOTS(1:5) .EQ. 'W-181') .OR. (MOTS(1:5) .EQ. 'W-185') .OR. (MOTS(1:5) .EQ. 'W-188')) THEN
        AF=0.025
*****
*
*   Alkaline earths
*

```

SPEFF.FOR

10-15-1987

```

*****
      ELSE IF (MOTS(1:2) .EQ. 'SR' .OR. (MOTS(1:2) .EQ. 'CA') .OR
&. (MOTS(1:2) .EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'BA')) THEN
      IF (PLIFE .GT. 15) THEN
      AF=0.025
      ELSE
      IF (EELEC(1) .LT. 0.2) THEN
      AF=0.25
      ELSE
      AF=0.025
      END IF
      END IF
      ELSE
*****
*      Electron emitter on bone surfaces      *
*****
      IF (EELEC(1) .LT. 0.2) THEN
      AF=0.25
      ELSE
      AF=0.025
      END IF
      END IF
*****
*      Cortical bone      *
*****
      ELSE IF (LOOP .GT. 1) THEN
*****
*      Electron emitter uniform in volume      *
*****
      IF (MOTS(1:4) .EQ. 'P-33' .OR. (MOTS(1:6) .EQ. 'NB-93M') .OR
&. (MOTS(1:5) .EQ. 'NB-94') .OR. (MOTS(1:5) .EQ. 'U-232') .OR. (MOTS(1:5) .EQ. 'U-233') .OR. (MOTS(1:5) .EQ. 'U-234') .OR. (MOTS(1:5) .EQ. 'U-235') .OR. (MOTS(1:5) .EQ. 'U-236') .OR. (MOTS(1:5) .EQ. 'U-238') .OR. (MOTS(1:2) .EQ. 'NA') .OR. (MOTS(1:2) .EQ. 'CR') .OR
&. (MOTS(1:2) .EQ. 'RB') .OR. (MOTS(1:5) .EQ. 'ZN-65') .OR. (MOTS(1:6) .EQ. 'PB-205') .OR. (MOTS(1:6) .EQ. 'PB-210') .OR. (MOTS(1:4) .EQ. 'BE-7') .OR. (MOTS(1:5) .EQ. 'BE-10') .OR. (MOTS(1:4) .EQ. 'V-49') .OR. (MOTS(1:6) .EQ. 'PD-103') .OR. (MOTS(1:6) .EQ. 'PD-107') .OR. (MOTS(1:6) .EQ. 'SN-113') .OR. (MOTS(1:7) .EQ. 'SN-119M') .OR. (MOTS(1:6) .EQ. 'SN-123') .OR. (MOTS(1:6) .EQ. 'SN-126') .OR. (MOTS(1:6) .EQ. 'TA-182') .OR. (MOTS(1:5) .EQ. 'W-181') .OR. (MOTS(1:5) .EQ. 'W-185') .OR. (MOTS(1:5) .EQ. 'W-186')) THEN
      AF=0.015
*****
*      Alkaline earths      *
*****
      ELSE IF (MOTS(1:2) .EQ. 'SR' .OR. (MOTS(1:2) .EQ. 'CA') .OR
&. (MOTS(1:2) .EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'BA')) THEN
      IF (PLIFE .GT. 15) THEN
      AF=0.015
      ELSE
      IF (EELEC(1) .LT. 0.2) THEN

```

SPEFF.FOR

10-15-1987

```

      AF=0.25
    ELSE
      AF=0.015
    END IF
  END IF
ELSE
*****
*      Electron emitter on bone surfaces      *
*****
      IF (EELEC(1) .LT. 0.2) THEN
        AF=0.25
      ELSE
        AF=0.015
      END IF
    END IF
  END IF
  SPEFF=SPEFF*((YELEC(1)*EELEC(1)*QF*AF)/120.)
*****
*
*      Red marrow
*
*****
  ELSE
*****
*      Trabecular bone
*
*****
  IF (LOOP .EQ. 1) THEN
*****
*      Electron emitter uniform in volume
*****
      IF (MOTS(1:4) .EQ. 'P-33' .OR. (MOTS(1:6) .EQ. 'NB-93M') .D
&R. (MOTS(1:5) .EQ. 'NB-94') .OR. (MOTS(1:5) .EQ. 'U-232') .OR. (M
&TS(1:5) .EQ. 'U-233') .OR. (MOTS(1:5) .EQ. 'U-234') .OR. (MOTS(1:5
&) .EQ. 'U-235') .OR. (MOTS(1:5) .EQ. 'U-236') .OR. (MOTS(1:5) .EQ.
& 'U-238') .OR. (MOTS(1:2) .EQ. 'NA') .OR. (MOTS(1:2) .EQ. 'CR') .D
&R. (MOTS(1:2) .EQ. 'RB') .OR. (MOTS(1:5) .EQ. 'ZN-65') .OR. (MOTS(
&1:6) .EQ. 'PB-205') .OR. (MOTS(1:6) .EQ. 'PB-210') .OR. (MOTS(1:4)
& .EQ. 'BE-7') .OR. (MOTS(1:5) .EQ. 'BE-10') .OR. (MOTS(1:4) .EQ. '
&V-49') .OR. (MOTS(1:6) .EQ. 'PD-103') .OR. (MOTS(1:6) .EQ. 'PD-107
&') .OR. (MOTS(1:6) .EQ. 'SN-113') .OR. (MOTS(1:7) .EQ. 'SN-119M')
&.OR. (MOTS(1:6) .EQ. 'SN-123') .OR. (MOTS(1:6) .EQ. 'SN-126') .OR.
& (MOTS(1:6) .EQ. 'TA-182') .OR. (MOTS(1:5) .EQ. 'W-181') .OR. (MOT
&S(1:5) .EQ. 'W-185') .OR. (MOTS(1:5) .EQ. 'W-188')) THEN
        AF=0.35
*****
*      Alkaline earths
*
*****
      ELSE IF (MOTS(1:2) .EQ. 'SR' .OR. (MOTS(1:2) .EQ. 'CA') .OR
&. (MOTS(1:2) .EQ. 'RA') .OR. (MOTS(1:2) .EQ. 'BA')) THEN
        IF (PLIFE .GT. 15) THEN
          AF=0.35
        ELSE

```

SPEFF.FOR

10-15-1987

```

      AF=0.5
      END IF
      ELSE
*****
*      Electron emitter on bone surfaces      *
*****
      AF=0.5
      END IF
      SPEFF=SPEFF+((YELEC(I)*EELEC(I)*QF*AF)/1500.)
*****
*      Cortical bone                          *
*****
      ELSE IF (LOOP .GT. 1) THEN
      AF=0.0
      END IF
      END IF
      END IF
180 CONTINUE
182 IF (LOOP .EQ. 0) GOTO 185
      GOTO 195
*****
*                                          *
*      Photon                          *
*                                          *
*****
185 IF (17 .EQ. 0) GOTO 195
      QF=1
      DO 190 I=1,17
      AF=0.
      IF (EGAMMA(I) .LT. 0.01) THEN
      IF (ICON .EQ. 1) THEN
      IF (JSORCE .EQ. 17) THEN
      AF=1.0/69900.
      SPEFF=SPEFF+(YGAMMA(I)*EGAMMA(I)*QF*AF)
      ELSE
      AF=0.
      END IF
      ELSE
      AF=1.0/THASS
      SPEFF=SPEFF+(YGAMMA(I)*EGAMMA(I)*QF*AF)
      END IF
      ELSE
      ICHECK=0
      AF=INTRPT(EGAMMA(I),JSORCE,KTARG,ICHECK)
      IF (ICHECK .EQ. 1) THEN
      WORD(1:5)='SORRY'
      RETURN
      END IF
      SPEFF=SPEFF+((YGAMMA(I)*EGAMMA(I)*QF*AF)/THASS)
      END IF
190 CONTINUE
195 SAVE=WORD

```

SPEFF.FOR

10-15-1987

RETURN
END

SUBMER.FOR

10-15-1987

```

*****
*
*      SUBROUTINE NAME : SUBMER FORTRAN
*      PURPOSE: Call data files "LIST" and "NOBLE" for dose
*               equivalent rate in target organs from eubereion
*               in e eemi-infinite cloud of noble gases or
*               elemental tritium
*
*      DESCRIPTION OF VARIABLES
*      -----
*      WORD ---> Name of the given isotope
*      NRATE ---> Dose equivalent rate in target organs or tissues
*                from eubereion in unit concentration of the
*                isotope
*      DER ----> Derived Air Concentration
*      RISK ---> DAC determined by the non-stochastic limit
*      ORGAN --> Tissue or organ of non-stochastic limit
*
*****
      SUBROUTINE SUBMER (WORD,NRATE,DER,RISK,ORGAN,*)
      DIMENSION NRATE(1:24),JSORCE(1:13),VALUE(1:13)
      CHARACTER*8 ERT,WORD
      OPEN(UNIT=10,FILE='LIST',FORM='FORMATTED',ACCESS='DIRECT',RECL=10)
      OPEN(UNIT=11,FILE='NOBLE',FORM='FORMATTED',ACCESS='DIRECT',RECL=16
&1)
      DD 2 I=1,24
      NRATE(I)=0.
2  CONTINUE
      IF (WORD(1:3) .EQ. 'H-3') THEN
          NRATE(1)=9.9E-15
          DER=2.0E10
          RISK=0.0E+00
          ORGAN=0
          RETURN
      END IF
      ITREK=0
      DD 5 LIS=1,26
      READ(10,6,REC=LIS,ERR=20)ERT,ICONT
6  FORMAT(A8,I2)
      IF (WORD .EQ. ERT) THEN
          ITREK=ICONT
          GOTD 7
      END IF
5  CONTINUE
      IF (ITREK .EQ. 0) GOTD 20
7  READ(11,9,REC=ITREK,ERR=25)DER,RISK,ORGAN,(JSORCE(I),VALUE(I),I=1,
&13)
9  FORMAT(1PEB.1,1PEB.1,13,12,1PEB.1,13,1PEB.1,13,1PEB.1,13,1PEB.1,13
&1,1PEB.1,13,1PEB.1,13,1PEB.1,13,1PEB.1,13,1PEB.1,13,1PEB.1,13,1PEB.
&1,13,1PEB.1,13,1PEB.1)
      DD 10 I=1,13
      IF (JSORCE(I) .EQ. 0) GOTD 10
      NRATE(JSORCE(I))=VALUE(I)
10 CONTINUE

```

SUBMER.FOR

10-15-1987

```
      RETURN
20  WRITE(*,21)
21  FORMAT(///,' Error: Nuclide not found in the ICRP Publication 30',/
&,' SOURCE: SUBMER FORTRAN',/,', CORRECTIVE ACTION: Try another nucl
&idel',////)
      RETURN 1
25  WRITE(*,26)
26  FORMAT(///,' ERROR: Unable to read dose equivalent rates from date
&file "NOBLE"! ',/,', SOURCE: SUBMER FORTRAN',/,', CORRECTIVE ACTION:
&check the file and try agein!',//)
      RETURN 1
END
```


TFRAC.FOR

11-02-1987

```

*****
*
*      SUBROUTINE NAME: TFRAC FORTRAN
*
*      PURPOSE: Retrieve the retention fractions, F2 and the
*               biological half-lives, BHALF of the nuclide
*               in source organs
*
*      DATA FILE REQUIRED: a) RETENT FILE
*
*      AUXILIARY SUBROUTINES REQUIRED:
*
*          a) I1 FORTRAN
*          b) SOURCE FORTRAN
*
*****

SUBROUTINE TFRAC(KZ,F2,BHALF,JSOURCE,SMASS,*)
  DIMENSION F2(1:3),BHALF(1:3)
  CHARACTER*20 C2
  OPEN (UNIT=15,FILE='RETENT',ACCESS='DIRECT',FORM='FORMATTED',RECL=
&66,STATUS='OLD')
  IF (KZ .GT. 92) THEN
    KZ=89
  END IF
  ISAVE=(5*(KZ-1))
  SUM=0.
  DO 1 I=1,5
    KEY=ISAVE+I
    READ(UNIT=15,FMT='(A20,F7.4,F7.4,F7.4,F7.4,F9.2,F8.1,F8.1)',ERR=10,REC=
&KEY)C2,D,E,F,G,H,B
*****
*
*      Function subprogram I1 converts C2, the source organ in
*      alphanumeric characters to an integer from source list to
*      compare with JSOURCE
*
*****

  IF (I .EQ. 1) THEN
    IF (D .EQ. 0. .AND. (G .EQ. 0.)) GOTO 10
  END IF
  IF (D .EQ. 0. .AND. (G .EQ. 0.)) GOTO 5
  ICOMP=I1(C2)
  IF (ICOMP .EQ. JSOURCE) THEN
    F2(1)=D
    F2(2)=E
    F2(3)=F
    BHALF(1)=G
    BHALF(2)=H
    BHALF(3)=B
*****
*
*      Function subprogram SOURCE gives the mass of the source
*      organ, when ICOMP is given as input
*
*****

  SMASS=SOURCE(ICOMP)
  GOTO 5
END IF

```

TFRAC.FOR

11-02-1987

```
*****
*
*   When source organ is 'Total body' and several organs of
*   mass Mi are associated with different retention fractions,
*   then the masses of all these organs are first summed, i.e.
*   (sum of Mi) and then 'Total body' is assigned a mass of
*   70000-(sum of Mi) and retention fractions which are
*   associated with 'all other'. The records are entered in
*   in such a way that for a given KZ, entry of 'all other' is
*   always at the end after all organs, i
*
*****
      IF (JSOURCE .EQ. 17) THEN
        IF (ICOMP .NE. 18) THEN
          SUM=SUM+SOURCE(ICOMP)
        ELSE
          SMASS=70000.-SUM
          FZ(1)=D
          FZ(2)=E
          FZ(3)=F
          BNALF(1)=G
          BNALF(2)=H
          BNALF(3)=B
        END IF
      END IF
      1 CONTINUE
      5 RETURN
      10 CALL CLEAR
          WRITE (*,15)
      15 FORMAT(/,' ERROR: Unable to read retention fractions, and biologi
&cal half-lives in source organ, from file "RETENT",/, ' SOURCE: T
&FRAC FORTRAN',/, ' CORRECTIVE ACTION: Try another nuclide!',////)
          PAUSE ' TO RESUME PRESS <RETURN>!!'
          RETURN 1
      END
```

THALF.FOR

11-02-1987

```
*****
*
*      FUNCTION SUBPROGRAM NAME: THALF FORTRAN
*
*      PURPOSE: Provide half-life of clearance from body fluid
*               compartment given an atomic number of a nuclide
*
*****
FUNCTION THALF(KZ)
  IF (KZ .EQ. 9 .OR. (KZ .EQ. 19) .OR. (KZ .EQ. 79) .OR. (KZ .EQ. 81
&))THEN
    THALF=0.
  ELSE IF (KZ .EQ. 15 .OR. (KZ .EQ. 24) .OR. (KZ .EQ. 27) .OR. (KZ .E
&Q. 90))THEN
    THALF=0.5
  ELSE IF (KZ .EQ. 43 .OR. KZ .EQ. 75)THEN
    THALF=0.02
  ELSE IF (KZ .EQ. 44 .OR. KZ .EQ. 45)THEN
    THALF=0.3
  ELSE IF (KZ .EQ. 52)THEN
    THALF=0.8
  ELSE IF (KZ .EQ. 83)THEN
    THALF=0.01
  ELSE
    THALF=0.25
  END IF
  RETURN
END
```

TRNSFM.FOR

11-02-1987

```

*****
*
*      SUBROUTINE NAME : TRNSFM FORTRAN
*      PURPOSE: Calculates source-organ transformations, US
*
*      DESCRIPTION OF VARIABLES
*      -----
*      UTJ(I)-----> Source-organ transformations of the isotope
*                      under consideration, j in the body fluid
*                      (transfer) compartment.
*      UROB(I)-----> The total number of transformations of
*                      radionuclide, i in 'rest of the body' of
*                      mass, 70000-(sum of Mj) where Mj is the
*                      mass of organ j for each unique retention
*                      fraction.
*      US(JSORCE,I)---> Source-organ transformations of nuclide, i
*      UJ-----> Transformations in each compartment of the
*                      source organ.
*
*****
      SUBROUTINE TRNSFM (FT,F2,BHALF,RCONST,NO,BRA,US,TCNST,JSORCE,F1,I
&PROG,FGI,SMAS,UROB,MROB,KZ)
      DIMENSION F2(1:3),BHALF(1:3),BCONST(1:3),RCONST(1:NO),BRA(1:NO),AS
&T(1:50),ASI(1:50),AULI(1:50),ALLI(1:50),US(1:20,1:NO),FT(1:NO),UTJ
&(1:50),FGI(1:NO),TEMP(1:3,1:50),UROB(1:NO)
      REAL MROB
*****
*
*      Calculating the biological constants,BCONST from the
*      biological half-lives in different compartments of the
*      source organ
*
*****
      DO 10 I=1,3
      IF (BHALF(I) .EQ. 0. .OR. BHALF(I) .EQ. 99999.9) THEN
        BCONST(I)=0.
      ELSE
        BCONST(I)=(LOG(2.)/BHALF(I))
      END IF
      10 CONTINUE
*****
*
*      There are separate methods of calculating source-organ
*      transformations for stomach, SI, LLI, ULI depending on the
*      mode of intake
*
*****
      IF (IPROG .EQ. 1) THEN
        IF (JSORCE .EQ. 2) GOTO 140
        IF (JSORCE .EQ. 3) GOTO 150
        IF (JSORCE .EQ. 4) GOTO 160
        IF (JSORCE .EQ. 5) GOTO 170
      ELSE IF (IPROG .EQ. 0) THEN
        IF (JSORCE .EQ. 2) GOTO 100

```

TRNSFM. FOR

11-02-1987

```

      IF (JSORCE .EQ. 3)GOTO 110
      IF (JSORCE .EQ. 4)GOTO 120
      IF (JSORCE .EQ. 5)GOTO 130
END IF
=====
*      Special treatment for iodine
=====
      IF (KZ .EQ. 53)GOTO 300
=====
*
*      Outer most loop for calculation of US of nuclide species,j
=====
      DO 15 J=1,N0
=====
*
*      In case of instantaneous transfer to tissue compartment,
*      calculation of UTJ(I) in transfer compartment is skipped
=====
      IF (TCONST .EQ. 0.)THEN
        DO 16 I=1,N0
          UTJ(I)=0.
16      CONTINUE
          GOTO 70
        END IF
=====
*
*      TRANSFER COMPARTMENT
=====
*
*      Loop for outer sum term,SUM in the calculation of UTJ(J)
=====
      SUM=0.
      DO 20 I=1,J
=====
*
*      Loop for outer product term,PROD of RCONST in the equation
*      for calculation of UTJ(J)
=====
      PROD=1.
=====
*
*      Parent case
=====
      IF (I+1 .GT. J)GOTO 30
=====
*
*      Daughters' case
=====
      DO 25 K=I+1,J
        PROD=FLOW(RCONST(K)*PROD)
25      CONTINUE
      SUM1=0.

```

TRNSFM. FOR

11-02-1987

```

*
*   Loop for inner sum term, SUM1 in the equation for
*   calculation of UTJ(J)
*
*****
DO 35 M=1,J
  PROD1=1.
*****
*
*   Loop for inner product term, PROD1 in the denominator in
*   equation for calculation of UTJ
*
*****
DO 40 K=1,J
  IF (K .EQ. M) GOTO 40
  PROD1=FLOW(PROD1*(RCONST(K)-RCONST(M)))
40 CONTINUE
  SUM1=SUM1+((1-UXP(-(TCONST+RCONST(M))*365.25*50.))/FLOW((TCONST+RCONST(M))*PROD1))
35 CONTINUE
  SUM=SUM+(PROD*FT(I))*SUM1/BRA(I)
20 CONTINUE
  UTJ(J)=SUM*BRA(J)
70 UJ=0.
*****
*
*   Loop for calculation of total source-organ transformations
*   from contribution of different compartments of each source
*
*****
DO 65 L=1,3
  IF (F2(L) .EQ. 0.) GOTO 65
  SUM2=0.
*****
*
*   TISSUE COMPARTMENT
*
*   Loop for outer sum term, SUM2 in calculation of source-organ
*   transformations, UJ in each compartment of the source organ
*
*****
DO 75 I=1,J
*****
*
*   Loop for outer product term, PROD2 in the equation for
*   calculation of UJ
*
*****
  PROD2=1.
*****
*
*   Parent case
*
*****
  IF (I+1 .GT. J) GOTO 85
*****

```

TRNSFM.FOR

11-02-1987

```

*                               Daughtere' case                               *
*****
      DO 80 K=1,J
        PROCD2=FLOW(RCONST(K)*PROCD2)
      80 CONTINUE
      85 SUM3=0.
*****
*      Loop for inner sum term, SUM3 in the equation      *
*****
      DO 90 M=1,J
        PROCD3=1.
*****
*      Loop for inner product term, PROCD3 in the equation  *
*****
      DO 95 K=1,J
        IF (K .EQ. M) GOTO 95
        PROCD3=FLOW(PROCD3*(RCONST(K)-RCONST(M)))
      95 CONTINUE
      SUM3=SUM3+(((1-LXP(-(BCONST(L)+RCONST(M))*365./2550))/FLOW((BCONST(
&L)+RCONST(M))*PROCD3))
      90 CONTINUE
*****
*      Initia! activity, ACTJ of species, j in source organ.      *
*****
      IF (TCONST .EQ. 0.) THEN
*****
*      Instantaneous transfer from body fluid compartment      *
*****
        ACTJ=F2(L)*FT(1)/BRA(1)
      ELSE
        ACTJ=FLOW(F2(L)*TCONST*UTJ(1)/BRA(1))
      END IF
      SUM2=SUM2+FLOW(PROCD2*ACTJ*SUM3)
      75 CONTINUE
      UJ=UJ+(BRA(J)*SUM2)
      65 CONTINUE
      US(JSORCE,J)=(SHASS*UTJ(J)/70000.)*UJ
      IF (JSORCE .EQ. 17) THEN
*****
*
*      Redistribute transformations uniformly through out all
*      organs and tissues of the body
*
*****
        UROB(J)=US(JSORCE,J)
        MROB=SHASS
*****
*      Source-organ transformations with source as 'total body'
*****
        US(JSORCE,J)=US(JSORCE,J)*70000./SHASS
      ELSE
        IF (UROB(J) .EQ. 0.) GOTO 15
*****
*      Reduce US(JSORCE,J) to compensate for redistribution of UROB(J)

```

TRNSFM. FOR

11-02-1987

```

*****
      US(JSORCE,J)=US(JSORCE,J)-(SMASS*UROB(J)/HROB)
      END IF
      15 CONTINUE
      GOTO 180
*****
      *          STOMACH in case of INGESTION          *
*****
      100 DO 105 J=1,NO
*****
      *          Parent case          *
*****
      IF (J .EQ. 1) THEN
        US(JSORCE,1)=BRA(1)/FLOW(24.*RCONST(1))
      ELSE
*****
      *          Daughters' case          *
*****
        US(JSORCE,J)=((US(JSORCE,J-1)/BRA(J-1))*RCONST(J)/FLOW(24.*RCONST(
        &J)))*BRA(J)
      END IF
      105 CONTINUE
      GOTO 180
*****
      *
      *          SMALL INTESTINE in case of INGESTION          *
      *
      *          If F1=1, it is assumed that the radionuclide passes directly
      *          from the stomach to body fluids and does not pass through
      *          other sections of the GI tract
      *
*****
      110 IF (F1 .EQ. 1.) GOTO 180
      BFCNST=F1*6./(1.-F1)
      DO 115 J=1,NO
*****
      *          Parent case          *
*****
      IF (J .EQ. 1) THEN
        US(JSORCE,1)=BRA(1)*24./FLOW((24.*RCONST(1))*6.+BFCNST+RCONST(1))
      ELSE
        AST(1)=1./FLOW(24.*RCONST(1))
      ELSE
*****
      *          Daughters' case          *
*****
        AST(J)=AST(J-1)*RCONST(J)/FLOW(24.*RCONST(J))
        R1=AST(J-1)*RCONST(J)*24./FLOW((24.*RCONST(J))*6.+BFCNST+RCONST(J)
        &J))
        R2=(US(JSORCE,J-1)/BRA(J-1))*RCONST(J)/FLOW(6.+BFCNST+RCONST(J))
        US(JSORCE,J)=BRA(J)*FLOW(R1+R2)
      END IF
      115 CONTINUE
      GOTO 180

```


TRNSFM. FOR

11-02-1987

```

*****
*      UPPER LARGE INTESTINE in case of INGESTION      *
*****
120 IF (F1 .EQ. 1.)GOTO 180
BFCNST=F1*6./(1.-F1)
DO 125 J=1,ND
*****
*      Parent case      *
*****
IF (J .EQ. 1)THEN
US(JSORCE,1)=BRA(J)*24.*6./FLOW((24.+RCONST(1))*(6.+BFCNST+RCONST(
&1)))*(1.8+RCONST(1)))
AST(1)=1./FLOW(24.+RCONST(1))
ASI(1)=24./FLOW((24.+RCONST(1))*(6.+BFCNST+RCONST(1)))
ELSE
*****
*      Daughters' case      *
*****
AST(J)=AST(J-1)*RCONST(J)/FLOW(24.+RCONST(J))
R1=AST(J-1)*RCONST(J)*24./FLOW((24.+RCONST(J))*(6.+BFCNST+RCONST(J
&)))
R2=ASI(J-1)*RCONST(J)/FLOW(6.+BFCNST+RCONST(J))
ASI(J)=FLOW(R1+R2)
R3=AST(J-1)*24.*6.*RCONST(J)/FLOW((24.+RCONST(J))*(6.+BFCNST+RCONS
&T(J))*(1.8+RCONST(J)))
R4=ASI(J-1)*RCONST(J)*6./FLOW((6.+BFCNST+RCONST(J))*(1.8+RCONST(J)
&))
R5=(US(JSORCE,J-1)/BRA(J-1))*RCONST(J)/FLOW(1.8+RCONST(J))
US(JSORCE,J)=FLOW(R3+R4+R5)*BRA(J)
END IF
125 CONTINUE
GOTO 180
*****
*      LOWER LARGE INTESTINE in case of INGESTION      *
*****
130 IF (F1 .EQ. 1.)GOTO 180
BFCNST=F1*6./(1.-F1)
DO 135 J=1,ND
R1=FLOW(24.+RCONST(J))
R2=FLOW(6.+BFCNST+RCONST(J))
R3=FLOW(1.8+RCONST(J))
R4=FLOW(1.+RCONST(J))
*****
*      Parent case      *
*****
IF (J .EQ. 1)THEN
US(JSORCE,1)=BRA(1)*24.*6.*1.8/FLOW(R1*R2*R3*R4)
AST(1)=1./R1
ASI(1)=24./FLOW(R1*R2)
AULI(1)=24.*6./FLOW(R1*R2*R3)
ELSE
*****
*      Daughters' case      *
*****

```

TRNSFM. FOR

11-02-1987

```

AST(J)=AST(J-1)*RCONST(J)/R1
ASI(J)=FLOW((AST(J-1)*RCONST(J)*24./(R1*R2))+ (ASI(J-1)*RCONST(J)/R
&2))
AULI(J)=FLOW((AST(J-1)*RCONST(J)*24.*(R1*R2*R3))+ (ASI(J-1)*RCON
&ST(J)*6./(R2*R3))+ (AULI(J-1)*RCONST(J)/(R3)))
US(JSORCE,J)=BRA(J)*FLOW((AST(J-1)*RCONST(J)*24.*1.8/(R1*R2*R3*
&R4))+ (ASI(J-1)*RCONST(J)*6.*1.8/(R2*R3*R4))+ (AULI(J-1)*RCONST(J)*1
&.8/(R3*R4))+ (US(JSORCE,J-1)/BRA(J-1))*RCONST(J)/R4))
END IF
135 CONTINUE
GOTO 180
*****
"          STOMACH in case of INHALATION          "
*****
140 DO 142 J=1,N0
*****
"          Parent case          "
*****
IF (J .EQ. 1) THEN
US(JSORCE,1)=BRA(1)*FGI(1)/FLOW(24.*RCONST(1))
*****
"          Daughtere' case          "
*****
ELSE
*****
"          Loop for outer sum term,SUM4          "
*****
SUM4=0.
DO 143 I=1,J
*****
"          Loop for outer product term,PROD4          "
*****
PROD4=1.
IF (I+1 .GT. J) GOTO 144
DO 145 K=I+1,J
PROD4=FLOW(RCONST(K)*PROD4)
145 CONTINUE
144 SUM5=0.
*****
"          Loop for inner sum term,SUM5          "
*****
DO 146 M=1,J
PROD5=1.
*****
"          Loop for inner product term,PROD5          "
*****
DO 147 N=1,J
IF (K .EQ. M) GOTO 147
PROD5=FLOW(PROD5*(RCONST(K)-RCONST(M)))
147 CONTINUE
SUM5=SUM5+((1-UXP(-(24.*RCONST(M))*365.25*50.))/FLOW((24.*RCONST(M
&2))*PROD5))
146 CONTINUE
SUM4=SUM4+FLOW(PROD4*FGI(1)*SUM5/BRA(I))

```

TRNSFM. FOR

11-02-1987

```

143 CONTINUE
    US(JSORCE,J)=SUM4*BRA(J)
    END IF
142 CONTINUE
    GOTO 180
*****
*          SMALL INTESTINE in case of INHALATION          *
*****
150 IF (F1 .EQ. 1.)GOTO 180
    BFCNST=F1*6./(1.-F1)
    OO 152 J=1,NO
*****
*          Parent case          *
*****
    IF (J .EQ. 1)THEN
        US(JSORCE,1)=BRA(1)*FGI(1)*24./FLOW((24.+RCONST(1)))*(6.+BFCNST+RCD
&NST(1)))
        INN=1
        DO 2 NUC=1,NO
            IF (NUC .EQ. 1)THEN
                TEMP(INN,NUC)=BRA(1)*FGI(1)/FLOW(24.+RCONST(1))
            ELSE
                *
                SUM4=0.
                DO 3 I=1,NUC
                    *****
                *          Loop for outer product term,PROD4          *
                *****
                PROD4=1.
                IF (I+1 .GT. NUC)GOTO 4
                DO 5 K=I+1,NUC
                    PROD4=FLOW(RCONST(K)*PROD4)
                5 CONTINUE
                4 SUM5=0.
                *****
                *          Loop for inner sum term,SUM5          *
                *****
                DO 6 N=I,NUC
                    PROD5=1.
                    *****
                *          Loop for inner product term,PROD5          *
                *****
                DO 7 K=I,NUC
                    IF (K .EQ. N)GOTO 7
                    PROD5=FLOW(PROD5*(RCONST(K)-RCONST(N)))
                7 CONTINUE
                SUM5=SUM5+((1.-UNP*(-(24.0+RCONST(N)))*365.25*50.))/FLOW((24.+RCONST
&(N))*PROD5))
                *
            6 CONTINUE
            SUM4=SUM4+FLOW(PROD4*FGI(1)*SUM5/BRA(1))
        3 CONTINUE
        TEMP(INN,NUC)=SUM4*BRA(NUC)
    END IF

```

TRNSFM.FOR

11-02-1987

```

2 CONTINUE
*****
*           Daughters' case           *
*****
ELSE
*****
*   Loop for outer sum term,SUM4      *
*****
    SUM4=0.
    DO 153 I=1,J
*****
*   Loop for outer product term,PROD4 *
*****
    PROD4=1.
    IF (I+1 .GT. J)GOTO 154
    DO 155 K=I+1,J
        PROD4=FLOW(RCONST(K)*PROD4)
155 CONTINUE
154 SUM5=0.
*****
*   Loop for inner sum term,SUM5      *
*****
    DO 156 M=1,J
        PROD5=1.
*****
*   Loop for inner product term,PROD5 *
*****
        DO 157 K=1,J
            IF (K .EQ. M)GOTO 157
            PROD5=FLOW(PROD5*(RCONST(K)-RCONST(M)))
157 CONTINUE
            SUM5=SUM5+((1.-EXP(-(6.*BFCNST+RCONST(M))*365.25*50.))/FLOW((6.*BFCNST+RCONST(M))*PROD5))
156 CONTINUE
        SUM4=SUM4+FLOW(PROD4*24.*TEMP(1,1)*SUM5/BRA(1))
153 CONTINUE
    US(JSORCE,J)=SUM4*BRA(J)
    END IF
152 CONTINUE
    GOTO 180
*****
*           UPPER LARGE INTESTINE in case of INHALATION           *
*****
160 IF (F1 .EQ. 1.)GOTO 180
    BFCNST=F1*6./(1.-F1)
    DO 162 J=1,NO
*****
*           Parent case                                           *
*****
    IF (J .EQ. 1)THEN
        US(JSORCE,1)=BRA(J)*FGI(J)*24.*6./FLOW((24.*RCONST(1))*(6.*BFCNST+RCONST(1))*(1.8+RCONST(1)))
        DO 102 INN=1,2
            DO 102 NUC=1,NO

```

TRNSFM. FOR

11-02-1987

```

IF (INN .EQ. 1 .AND. (NUC .EQ. 1))THEN
  TEMP(INN,NUC)=BRA(1)*FGI(1)/FLOW(24.+RCONST(1))
ELSE IF (INN .EQ. 2 .AND. (NUC .EQ. 1))THEN
  TEMP(INN,NUC)=BRA(1)*FGI(1)*24./FLOW((24.+RCONST(1))*(6.+BFCNST
&+RCONST(1)))
ELSE
*
  SUM4=0.
  DO 103 I=1,NUC
*****
*   Loop for outer product term,PROD4
*****
    PROD4=1.
    IF (I+1 .GT. NUC)GOTO 104
    DO 108 K=I+1,NUC
      PROD4=FLOW(RCONST(K)*PROD4)
    108 CONTINUE
    104 SUM5=0.
*****
*   Loop for inner sum term,SUM5
*****
    DO 106 M=I,NUC
      PROD5=1.
*****
*   Loop for inner product term,PROD5
*****
      DO 107 K=I,NUC
        IF (K .EQ. M)GOTO 107
        PROD5=FLOW(PROD5*(RCONST(K)-RCONST(M)))
      107 CONTINUE
      IF (INN .EQ. 1)THEN
        SUM5=SUM5+((1.-UXP(-(24.0+RCONST(M))*365.25*50.))/FLOW((24.+RCON
&NST(M))*PROD5))
      ELSE IF (INN .EQ. 2)THEN
        SUM5=SUM5+((1.-UXP(-(6.+BFCNST+RCONST(M))*365.25*50.))/FLOW((6.
&+BFCNST+RCONST(M))*PROD5))
      END IF
*
    106 CONTINUE
    IF (INN .EQ. 1)THEN
      SUM4=SUM4+FLOW(PROD4*FGI(1))*SUM5/BRA(1))
    ELSE IF (INN .EQ. 2)THEN
      SUM4=SUM4+FLOW(PROD4*TEMP(1,I)*24.*SUM5/BRA(1))
    END IF
  103 CONTINUE
  TEMP(INN,NUC)=SUM4*BRA(NUC)
  END IF
  102 CONTINUE
*****
*   Daughters' case
*****
  ELSE
*****
*   Loop for outer sum term,SUM4
*****

```

TRNSEM.FOR

11-02-1987

```

*****
      SUM4=0.
      DO 163 I=1,J
*****
*      Loop for outer product term,PROD4
*****
      PROD4=1.
      IF (I+1 .GT. J)GOTO 164
      DO 165 K=I+1,J
      PROD4=FLOW(RCONST(K)*PROD4)
165 CONTINUE
164 SUM5=0.
*****
*      Loop for inner sum term,SUM5
*****
      DO 166 M=I,J
      PROD5=1.
*****
*      Loop for inner product term,PROD5
*****
      DO 167 K=I,J
      IF (K .EQ. M)GOTO 167
      PROD5=FLOW(PROD5*(RCONST(K)-RCONST(M)))
167 CONTINUE
      SUM5=SUM5+((1.-UXP(-(1.8+RCONST(M))*365.25*50.))/FLOW((1.8+RCONST(
      M))*PROD5))
166 CONTINUE
      SUM4=SUM4+FLOW(PROD4*6.*TEMP(2,I)*SUM5/BRA(1))
163 CONTINUE
      US(JSORCE,J)=SUM4*BRA(J)
      END IF
162 CONTINUE
      GOTO 180
*****
*      LOWER LARGE INTESTINE in case of INHALATION
*****
170 IF (F1 .EQ. 1.)GOTO 180
      BFCNST=F1*6./(1.-F1)
      DO 172 J=1,N0
      R1=FLOW(24.*RCONST(J))
      R2=FLOW(6.*BFCNST+RCONST(J))
      R3=FLOW(1.8+RCONST(J))
      R4=FLOW(1.*RCONST(J))
*****
*      Parent case
*****
      IF (J .EQ. 1)THEN
      US(JSORCE,1)=BRA(1)*FGI(J)*24.*6.*1.8/FLOW(R1*R2*R3*R4)
      DO 202 INN=1,3
      DO 202 NUC=1,N0
      IF (INN .EQ. 1 .AND. (NUC .EQ. 1))THEN
      TEMP(INN,NUC)=BRA(1)*FGI(1)/FLOW(24.*RCONST(1))
      ELSE IF (INN .EQ. 2 .AND. (NUC .EQ. 1))THEN
      TEMP(INN,NUC)=BRA(1)*FGI(1)*24./FLOW((24.*RCONST(1))*6.*BFCNST

```

TRNSFM. FOR

11-02-1987

```

&+RCONST(1)))
ELSE IF (INN .EQ. 3 .AND. (NUC .EQ. 1))THEN
    TEMP(INN,NUC)=BRA(1)*FGI(1)*24.*6./FLOW((24.*RCONST(1))*(6.*BFC
&NST+RCONST(1))*(1.8+RCONST(1)))
ELSE
*
    SUM4=0.
    DO 203 I=1,NUC
*****
*   Loop for outer product term,PROD4
*****
        PROD4=1.
        IF (I+1.GT. NUC)GOTO 204
        DO 205 K=I+1,NUC
            PROD4=FLOW(RCONST(K)*PROD4)
        205 CONTINUE
        204 SUM5=0.
*****
*   Loop for inner sum term,SUM5
*****
        DO 206 N=I,NUC
            PROD5=1.
*****
*   Loop for inner product term,PROD5
*****
            DO 207 K=I,NUC
                IF (K .EQ. N)GOTO 207
                PROD5=FLOW(PROD5*(RCONST(K)-RCONST(N)))
            207 CONTINUE
            IF (INN .EQ. 1)THEN
                SUM5=SUM5+((1.-UXP(-(24.0+RCONST(N))*365.25*50.))/FLOW((24.*ROD
&NST(N))*PROD5))
            ELSE IF (INN .EQ. 2)THEN
                SUM5=SUM5+((1.-UXP(-(6.*BFCNST+RCONST(N))*365.25*50.))/FLOW((6.
&0+BFCNST+RCONST(N))*PROD5))
            ELSE IF (INN .EQ. 3)THEN
                SUM5=SUM5+((1.-UXP(-(1.8+RCONST(N))*365.25*50.))/FLOW((1.8+RCON
&ST(N))*PROD5))
            END IF
*
        206 CONTINUE
        IF (INN .EQ. 1)THEN
            SUM4=SUM4+FLOW(PROD4*FGI(1))*SUM5/BRA(1))
        ELSE IF (INN .EQ. 2)THEN
            SUM4=SUM4+FLOW(PROD4*TEMP(1,1)*24.*SUM5/BRA(1))
        ELSE IF (INN .EQ. 3)THEN
            SUM4=SUM4+FLOW(PROD4*TEMP(2,1)*6.*SUM5/BRA(1))
        END IF
    203 CONTINUE
    TEMP(INN,NUC)=SUM4*BRA(NUC)
    END IF
    202 CONTINUE
*
*****

```

TRNSFM.FOR

11-02-1987

```

*                               Daughters' case                               *
*****
ELSE
*****
*   Loop for outer sum term, SUM4   *
*****
    SUM4=0.
    DO 173 I=1,J
*****
*   Loop for outer product term, PROD4   *
*****
    PROD4=1.
    IF (I+1 .GT. J) GOTO 174
    DO 175 K=I+1,J
    PROD4=FLOW(RCONST(K)*PROD4)
*****
175 CONTINUE
174 SUM5=0.
*****
*   Loop for inner sum term, SUM5   *
*****
    DO 176 M=I,J
    PROD5=1.
*****
*   Loop for inner product term, PROD5   *
*****
    DO 177 K=I,J
    IF (K .EQ. M) GOTO 177
    PROD5=FLOW(PROD5*(RCONST(K)-RCONST(M)))
177 CONTINUE
    SUM5=SUM5+((1.-EXP(-(1.0+RCONST(M))*365.25*50.))/FLOW((1.0+RCONST(
    &M))*PROD5))
176 CONTINUE
    SUM4=SUM4+FLOW(PROD4*1.8*TEMP(3,I)*SUM5/BRA(I))
173 CONTINUE
    US(JSORCE,J)=SUM4*BRA(J)
    END IF
172 CONTINUE
    GOTO 180
*****
*   Three compartment model for iodine   *
*****
300 DO 310 I=1,N0
    F2I=0.3
    F4I=0.9
    BODY=LOG(2.)/120
    BSORCE=LOG(2.)/12
    PART1=(RCONST(I)+TCONST)*(RCONST(I)+BODY)*(RCONST(I)+BSORCE)
    PART2=F2I*F4I*TCONST*BODY*BSORCE
    DENOM=PART1-PART2
    UBF=FT(I)*(RCONST(I)+BODY)*(RCONST(I)+BSORCE)/DENOM
    IF (JSORCE .EQ. 17) THEN
        UROB(I)=(F2I*TCONST*BODY*FT(I)/DENOM)*(SMASS*UBF/70000.)
        MROB=SMASS
        US(JSORCE,I)=UROB(I)*70000./SMASS
    
```


TRNSFM.FOR

11-02-1987

```
ELSE IF (JSORCE .EQ. 16) THEN
  US(JSORCE, I) = (F21 * TCONST * FT(I) * (RCONST(1) + BSORCE) / DENOM) + (SMASS *
&UBF / 70000.) - (SMASS * UROB(I) / WROB)
ELSE
  US(JSORCE, I) = 0.
END IF
310 CONTINUE
180 DO 500 I = 1, N0
  IF (US(JSORCE, I) .LT. 0.) THEN
    US(JSORCE, I) = 0.
  END IF
500 CONTINUE
RETURN
END
FUNCTION FLOW(ARGUM)
IF (ABS(ARGUM) .LT. 1.E-30) THEN
  FLOW = 1.E-30
ELSE IF (ABS(ARGUM) .GT. 1.E30) THEN
  FLOW = 1.E30
ELSE
  FLOW = ARGUM
END IF
RETURN
END
```

UXP.FOR

11-02-1987

```
FUNCTION UXP(ARG) .  
  IF (ARG .GT. -80) THEN  
    UXP = EXP(ARG)  
  ELSE  
    UXP = 0.  
  END IF  
  RETURN  
END
```

YERROR.FOR

11-02-1987

```
*****
*
*      SUBROUTINE NAME : YERROR FORTRAN
*      PURPOSE: Error handling subroutine for DECAY FORTRAN
*
*****
      SUBROUTINE YERROR(DECERR)
      INTEGER DECERR(7)
      IF (DECERR(1) .GT. 0) THEN
        WRITE (*,5)
5       FORMAT(//,' ERROR: No match found in file "ISOTIPS" for the gi
&ven radionuclide',/,,' SOURCE: DECAY FORTRAN',/,,' CORRECTIVE ACTID
&M: Try another nuclide !',/,/,/,/)
        DECERR(1)=0
        RETURN
      ELSE IF (DECERR(2) .GT. 0) THEN
        WRITE (*,10)
10      FORMAT(//,' ERROR: Unable to read the decay scheme of the nuc
&lide in file "ISOTOPE"',/,,' SOURCE: DECAY FORTRAN',/,,' CORRECTIVE
&ACTION: Try another nuclide!',/,/,/,/)
        DECERR(2)=0
        RETURN
      ELSE IF (DECERR(3) .GT. 0) THEN
        WRITE (*,15)
15      FORMAT(//,' ERROR: Unable to read the elphe energy and intens
&ity from file "ALPHA" for the nuclide',/,,' SOURCE: DECAY FORTRAN
&',/,,' CORRECTIVE ACTION: Try another nuclide!',/,/,/,/)
        DECERR(3)=0
        RETURN
      ELSE IF (DECERR(4) .GT. 0) THEN
        WRITE (*,20)
20      FORMAT(//,'ERROR: Unable to read the beta energy and intensit
&Y FROM FILE "BETA" FOR THE NUCLIDE',/,,' SOURCE: DECAY FORTRAN
&',/,,' CORRECTIVE ACTION: TRY ANOTHER NUCLIDE!',/,/,/,/)
        DECERR(4)=0
        RETURN
      ELSE IF (DECERR(5) .GT. 0) THEN
        WRITE (*,25)
25      FORMAT(//,'ERROR: Unable to read the positron energy and inten
&SITY FROM FILE "POSITRN" FOR THE NUCLIDE',/,,' SOURCE: DECAY FORT
&AN',/,,' CORRECTIVE ACTION: TRY ANOTHER NUCLIDE!',/,/,/,/)
        DECERR(5)=0
        RETURN
      ELSE IF (DECERR(6) .GT. 0) THEN
        WRITE (*,30)
30      FORMAT(//,'ERROR: Unable to read the electron energy end inten
&eity from file "ELECTRN" for the nuclide',/,,' SOURCE: DECAY FORT
&AN',/,,' CORRECTIVE ACTION: Try another nuclide!',/,/,/,/)
        DECERR(6)=0
        RETURN
      ELSE IF (DECERR(7) .GT. 0) THEN
        WRITE (*,35)
35      FORMAT(//,'ERROR: Unable to read the photon energy end intensi
&ty from file "PHOTON" for the nuclide',/,,' SOURCE: DECAY FORTRAN'
```

YERROR.FOR

11-02-1987

```
&,/, ' CORRECTIVE ACTION: Try another nuclide !!',////)  
  DECERR(7)=0  
  RETURN  
END IF  
END
```

A COMPUTER PROGRAM FOR INTERNAL DOSIMETRY ANALYSIS
USING THE METHODS OF ICRP-30

by

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ABSTRACT

A software package written in the FORTRAN-77 language uses the methods described in Part 1 of Publication 30 of the International Commission on Radiological Protection (ICRP) to calculate committed dose equivalents from an internal radionuclide to organs and tissues of an adult "reference man". An alternate version for use on an IBM-PC or a compatible micro-computer is also available.

The program considers any of the three major modes of intake of a radionuclide, namely, ingestion, inhalation, or submersion in a cloud of inert radioactive gas or elemental tritium. Except for the source of radiological decay data, the general principles, definitions, mathematical models, and calculational procedures follow closely those described in ICRP-30. The program calculates specific committed dose equivalents (Sv/Bq) in 19 target organs, annual limit of intake (Bq), and derived air concentration (in the case of inhalation or submersion) of a radionuclide. Also, weighted committed dose equivalents (Sv/Bq) for selected target organs receiving greater than or equal to 10 percent of the maximum dose is shown. In addition, a table of specific effective energies in 17 sources and 19 target organs in three units (MeV/g, rad/ μ Ci.h, and mSv/GBq.h), and a table for number of transformations in source organs per unit intake of activity of the radionuclide (/Bq) can be generated upon request. The features of independent subroutines and alterable data files on radiological decay and biological characteristics make modifications and update of the program simple and straightforward.